

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 13, 2004, 07:26:14 ; Search time 53 Seconds
(without alignments)
95.960 Million cell updates/sec

Title: US-09-747-029b-17

Perfect score: 105

Sequence: 1 QDTIVGWGDSXGCRPGQ 18

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 481716

Minimum DB seq length: 0

Maximum DB seq length: 18

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

- Database : A_Geneseq_29Jan04:*
- 1: Geneseqp1980s:*
 - 2: Geneseqp1990s:*
 - 3: Geneseqp2000s:*
 - 4: Geneseqp2001s:*
 - 5: Geneseqp2002s:*
 - 6: Geneseqp2003as:*
 - 7: Geneseqp2003Bs:*
 - 8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	103	98.1	18	4	Aae07230 IGP1685 p
2	83	79.0	14	4	Aae07231 IGP1686 p
3	60	57.1	18	4	Aae07225 IGP1650 p
4	47	44.8	16	4	Aab76381 Erbb2 bin
5	47	44.8	16	4	Aab76385 Erbb2 bin
6	44	41.9	14	4	Aae07229 IGP1684 p
7	44	41.9	16	4	Aab76382 Erbb2 bin
8	43	41.0	14	4	Aae07227 IGP1676 p
9	43	41.0	18	4	Aae07223 IGP1648 p
10	42	40.0	16	4	Aab76392 Erbb2 bin
11	42	40.0	16	4	Aab76387 Erbb2 bin
12	42	40.0	17	4	Aab76369 Erbb2 bin
13	42	40.0	17	4	Aab76370 Erbb2 bin
14	42	40.0	18	4	Aae07221 IGP1646 p
15	41	39.0	16	4	Aab76383 Erbb2 bin
16	41	39.0	16	4	Aab76390 Erbb2 bin
17	41	39.0	17	4	Aab76354 Erbb2 bin
18	40	38.1	14	4	Aae07228 IGP1687 p
19	39	37.1	11	2	Aaw05559 Thrombopo
20	39	37.1	11	2	Aaw36710 Thrombopo
21	39	37.1	11	4	Aau25929 Human thr
22	39	37.1	18	4	Aae07224 IGP1649 p
23	38	36.2	14	2	Aay08356 Cysteine
24	38	36.2	15	4	Aaj03265 Hepatitis
25	38	36.2	15	4	Aaj03570 Hepatitis

26	38	36.2	15	4	Aaj03655 Hepatitis
27	38	36.2	17	4	Aab76353 Erbb2 bin
28	38	36.2	17	4	Aab76373 Erbb2 bin
29	38	36.2	18	4	Aae07220 IGP1611 p
30	37	35.2	12	4	Aae06030 Dodecamer
31	37	35.2	15	2	Aar93669 HIV princ
32	37	35.2	16	4	Aab76384 Erbb2 bin
33	37	35.2	17	2	Aar06085 Immunorea
34	36	34.3	10	5	Abg98831 F protein
35	36	34.3	10	5	Abg98832 F protein
36	36	34.3	10	5	Abg98833 F protein
37	36	34.3	16	4	Aab76388 Erbb2 bin
38	36	34.3	16	4	Aab76386 Erbb2 bin
39	36	34.3	16	4	Aab76381 Erbb2 bin
40	36	34.3	16	4	Aag73166 Protease
41	36	34.3	17	4	Aam15636 Peptide #
42	36	34.3	17	4	Aab76372 Erbb2 bin
43	36	34.3	17	4	Aab76368 Erbb2 bin
44	36	34.3	17	4	Aab76371 Erbb2 bin
45	36	34.3	17	4	Aab76375 Erbb2 bin

ALIGNMENTS

RESULT 1
AAE07230
ID AAE07230 standard; peptide; 18 AA.

AC AAE07230;

XX 06-NOV-2001 (first entry)

DT IGP1685 peptide for diagnosis and treatment of rheumatoid arthritis.

DE Synthetic peptide; cyclic; IGP1685; autoimmune antibody;

XX rheumatoid arthritis; therapy; autoimmune disease; anti-rheumatic;

KW systemic hyporesponsiveness; immunosuppressive; antiarthritic.

OS Synthetic.

XX Key Location/Qualifiers

FT Modified-site 1..18 /note= "Biotinylated residues"

FT Disulfide-bond 9..14

FT Modified-site 12 /note= "Citrulline"

XX WO200146222-A2.

XX 28-JUN-2001.

XX 20-DEC-2000; 2000WO-EP013037.

XX 21-DEC-1999; 99EP-00870280.

XX 08-SEP-2000; 2000EP-00870195.

XX (INNO-) INNOGENETICS NV.

XX Union A, Moereels H, Meheus L;

XX WPI; 2001-496657/54.

XX New peptides, useful for diagnosing and treating rheumatoid arthritis, comprises citrulline residue between 2 cysteine residues and is specifically recognized by autoimmune antibodies from patients suffering from rheumatoid arthritis.

XX Claim 9; Page 42; 53pp; English.

XX The present sequence is a cyclic synthetic biotinylated peptide, IGP1685. The peptide comprises a citrulline residue between 2 cysteine residues and is specifically recognised by autoimmune antibodies from patients

CC suffering from rheumatoid arthritis. The peptide comprises amino acids
 CC involved in side chain interactions which is essential for the formation
 CC of three-dimensional structure of the peptide. The peptide of the
 CC invention is useful as a medicament to treat autoimmune diseases,
 CC preferably rheumatoid arthritis. It is also useful for treating
 CC autoimmune diseases by increasing the size of antigen-immune complexes to
 CC improve clearance of the formed immune complexes and for the preparation
 CC of a medicament for oral or nasal administration to treat autoimmune
 CC diseases by inducing a state of systemic hyporesponsiveness or tolerance
 CC to the peptide
 XX
 SQ Sequence 18 AA;

Query Match 98.1%; Score 103; DB 4; Length 18;
 Best Local Similarity 100.0%; Pred. No. 6.2e-07; Indels 0; Gaps 0;
 Matches 18; Conservative 0; Mismatches 0;

QY 1 QDTIVGWGCDXGCRPGQ 18
 |||||
 Db 1 QDTIVGWGCDXGCRPGQ 18
 |||||

RESULT 2
 AAE07231
 ID AAE07231 standard; peptide; 14 AA.
 AC
 XX AAE07231;
 XX

DT 06-NOV-2001 (first entry)

XX IGP1696 peptide for diagnosis and treatment of rheumatoid arthritis.

DE
 XX Synthetic peptide; cyclic; IGP1686; autoimmune antibody;
 KW rheumatoid arthritis; therapy; autoimmune disease; antirheumatic;
 KW systemic hyporesponsiveness; immunosuppressive; antiarthritic.
 XX
 OS Synthetic.

XX Key Location/Qualifiers
 FH Modified-site 1..14
 FT /note= "Biotinylated residues"
 FT Disulfide-bond 9..14
 FT Modified-site 12
 FT /note= "Citrulline"

XX WO200146222-A2.

XX 28-JUN-2001.

XX 20-DEC-2000; 2000WO-EP013037.

XX 21-DEC-1999; 99EP-00870280.

XX 08-SEP-2000; 2000EP-00870195.

XX (INNO-) INNOGENETICS NV.

XX Union A, Moereels H, Meheus L;

XX WPI; 2001-496657/54.

XX New peptides, useful for diagnosing and treating rheumatoid arthritis,
 PT comprises citrulline residue between 2 cysteine residues and is
 PT specifically recognized by autoimmune antibodies from patients suffering
 PT from rheumatoid arthritis.

XX Claim 9; Page 42; 53pp; English.

XX The present sequence is a cyclic synthetic biotinylated peptide, IGP1686.
 CC The peptide comprises a citrulline residue between 2 cysteine residues
 CC and is specifically recognised by autoimmune antibodies from patients
 CC suffering from rheumatoid arthritis. The peptide comprises amino acids
 CC involved in side chain interactions which is essential for the formation
 CC of three-dimensional structure of the peptide. The peptide of the

CC invention is useful as a medicament to treat autoimmune diseases,
 CC preferably rheumatoid arthritis. It is also useful for treating
 CC autoimmune diseases by increasing the size of antigen-immune complexes to
 CC improve clearance of the formed immune complexes and for the preparation
 CC of a medicament for oral or nasal administration to treat autoimmune
 CC diseases by inducing a state of systemic hyporesponsiveness or tolerance
 CC to the peptide
 XX
 SQ Sequence 14 AA;

Query Match 79.0%; Score 83; DB 4; Length 14;
 Best Local Similarity 100.0%; Pred. No. 0.00023;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 VGVGCDXGCRPGQ 18
 |||||
 Db 1 VGVGCDXGCRPGQ 14
 |||||

RESULT 3
 AAE07225

ID AAE07225 standard; peptide; 18 AA.

XX AC AAE07225;

XX DT 06-NOV-2001 (first entry)

XX IGP1650 peptide for diagnosis and treatment of rheumatoid arthritis.

XX Synthetic peptide; cyclic; IGP1650; autoimmune antibody;
 KW rheumatoid arthritis; therapy; autoimmune disease; antirheumatic;
 KW systemic hyporesponsiveness; immunosuppressive; antiarthritic.

XX OS Synthetic.

XX Key Location/Qualifiers
 FH Modified-site 1..18
 FT /note= "Biotinylated residues"
 FT Disulfide-bond 9..14
 FT Modified-site 11
 FT /note= "Citrulline"
 FT Modified-site 12
 FT /note= "Citrulline"

XX WO200146222-A2.

XX 28-JUN-2001.

XX 20-DEC-2000; 2000WO-EP013037.

XX 21-DEC-1999; 99EP-00870280.

XX 08-SEP-2000; 2000EP-00870195.

XX (INNO-) INNOGENETICS NV.

XX Union A, Moereels H, Meheus L;

XX WPI; 2001-496657/54.

XX New peptides, useful for diagnosing and treating rheumatoid arthritis,
 PT comprises citrulline residue between 2 cysteine residues and is
 PT specifically recognized by autoimmune antibodies from patients suffering
 PT from rheumatoid arthritis.

XX Claim 9; Page 42; 53pp; English.

XX The present sequence is a cyclic synthetic biotinylated peptide, IGP1650.
 CC The peptide comprises a citrulline residue between 2 cysteine residues
 CC and is specifically recognised by autoimmune antibodies from patients
 CC suffering from rheumatoid arthritis. The peptide comprises amino acids
 CC involved in side chain interactions which is essential for the formation
 CC of three-dimensional structure of the peptide. The peptide of the
 CC invention is useful as a medicament to treat autoimmune diseases,

CC preferably rheumatoid arthritis. It is also useful for treating
 CC autoimmune diseases by increasing the size of antigen-immune complexes to
 CC improve clearance of the formed immune complexes and for the preparation
 CC of a medicament for oral or nasal administration to treat autoimmune
 CC diseases by inducing a state of systemic hyporesponsiveness or tolerance
 CC to the peptide
 CC
 XX
 SQ Sequence 18 AA;

Query Match 57.1%; Score 60; DB 4; Length 18;
 Best Local Similarity 70.6%; Pred. No. 0.32;
 Matches 12; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 QDTIVGWGCDXSGCRPG 17
 |||||
 DB 1 QDTIHGFCXXGCRPG 17
 |||||

RESULT 4
 AAB76391
 ID AAB76391 standard; peptide; 16 AA.

AC AAB76391;

DT 10-APR-2001 (first entry)

DE Erbb2 binding peptide amino acid sequence SEQ ID 42.

KW Human; erbb2; HER2; cancer; nervous system disease; stroke; ischaemia;
 KW metabolic disorder; nutritional deficiency; Alzheimer's disease;
 KW Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
 KW Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.

OS Synthetic.

XX WO200101748-A2.

XX 11-JAN-2001.

PF 30-JUN-2000; 2000WO-US018283.

PR 02-JUL-1999; 99US-0142232P.

XX (GETH) GENENTECH INC.

XX Dennis MS;

XX WPI; 2001-123048/13.

XX Non-naturally occurring peptide ligands which compete for binding human
 PT erB2 gene products, useful for treating e.g. Alzheimer's disease,
 PT multiple sclerosis and diabetic neuropathy.

XX Disclosure; Fig 16; 116pp; English.

XX This invention relates to non-naturally occurring peptide ligands which
 CC bind to the human erB2 gene product Erbb2 (also known as HER2). Peptides
 CC represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
 CC of the Erbb2 binding ligands of the invention. Sequences AAB76421 -
 CC AAB76431 represent antibody Fc amino acid sequences used in the isolation
 CC of the peptides of the invention. The peptides compete for binding Erbb2
 CC with naturally occurring ligands, and may be used to treat disorders
 CC characterized by over expression of HER2/Erbb2 such as cancers, diseases
 CC of the nervous system, musculature and epithelia, e.g. nervous system
 CC damage resulting from trauma, surgery, strokes, ischaemia, infection,
 CC metabolic disorders, nutritional deficiency or toxic agents. In
 CC particular the synthetic peptide ligands may be used to treat Alzheimer's
 CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
 CC chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
 CC associated with diabetes

XX Sequence 16 AA;

Query Match 44.8%; Score 47; DB 4; Length 16;
 Best Local Similarity 70.0%; Pred. No. 15;
 Matches 7; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 6 GWGCDXSGCR 15
 |||||
 DB 5 GWGCIQPGCR 14
 |||||

RESULT 5
 AAB76385
 ID AAB76385 standard; peptide; 16 AA.

XX AAB76385;

DT 10-APR-2001 (first entry)

DE Erbb2 binding peptide amino acid sequence SEQ ID 36.

KW Human; erbb2; HER2; cancer; nervous system disease; stroke; ischaemia;
 KW metabolic disorder; nutritional deficiency; Alzheimer's disease;
 KW Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
 KW Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.

OS Synthetic.

XX WO200101748-A2.

XX 11-JAN-2001.

PF 30-JUN-2000; 2000WO-US018283.

PR 02-JUL-1999; 99US-0142232P.

XX (GETH) GENENTECH INC.

XX Dennis MS;

XX WPI; 2001-123048/13.

XX Non-naturally occurring peptide ligands which compete for binding human
 PT erB2 gene products, useful for treating e.g. Alzheimer's disease,
 PT multiple sclerosis and diabetic neuropathy.

XX Disclosure; Fig 16; 116pp; English.

XX This invention relates to non-naturally occurring peptide ligands which
 CC bind to the human erB2 gene product Erbb2 (also known as HER2). Peptides
 CC represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
 CC of the Erbb2 binding ligands of the invention. Sequences AAB76421 -
 CC AAB76431 represent antibody Fc amino acid sequences used in the isolation
 CC of the peptides of the invention. The peptides compete for binding Erbb2
 CC with naturally occurring ligands, and may be used to treat disorders
 CC characterized by over expression of HER2/Erbb2 such as cancers, diseases
 CC of the nervous system, musculature and epithelia, e.g. nervous system
 CC damage resulting from trauma, surgery, strokes, ischaemia, infection,
 CC metabolic disorders, nutritional deficiency or toxic agents. In
 CC particular the synthetic peptide ligands may be used to treat Alzheimer's
 CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
 CC chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
 CC associated with diabetes

XX Sequence 16 AA;

Query Match 44.8%; Score 47; DB 4; Length 16;
 Best Local Similarity 70.0%; Pred. No. 15;
 Matches 7; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 6 GWGCDXSGCR 15
 |||||
 DB 5 GWGCIQPGCR 14
 |||||

```

RESULT 6
AAE07229
ID AAE07229 standard; peptide; 14 AA.
XX
AC AAE07229;
XX
XX 06-NOV-2001 (first entry)
XX
DE IGP1684 peptide for diagnosis and treatment of rheumatoid arthritis.
XX
XX Synthetic peptide; cyclic; IGP1684; autoimmune antibody;
XX rheumatoid arthritis; therapy; autoimmune disease; anti-rheumatic;
XX systemic hyporesponsiveness; immunosuppressive; antiarthritic.
XX
OS Synthetic.
XX
XX Key Location/Qualifiers
FH Modified-site 1..14
FT /note= "Biotinylated residues"
FT Disulfide-bond 9..16
FT Modified-site 12
FT /note= "Citrulline"
XX
XX WO200146222-A2.
XX
XX 28-JUN-2001.
XX
XX 20-DEC-2000; 2000WO-EP013037.
XX
XX 21-DEC-1999; 99EP-00870280.
XX
XX 08-SEP-2000; 2000EP-00870195.
XX
XX (INNO-) INNOGENETICS NV.
XX
XX Union A, Moereels H, Meheus L;
XX
XX WPI; 2001-496657/54.
XX
XX New peptides, useful for diagnosing and treating rheumatoid arthritis,
XX comprises citrulline residue between 2 cysteine residues and is
XX specifically recognized by autoimmune antibodies from patients suffering
XX from rheumatoid arthritis.
XX
XX Claim 9; Page 42; 53pp; English.
XX
XX The present sequence is a cyclic synthetic biotinylated peptide, IGP1684.
XX The peptide comprises a citrulline residue between 2 cysteine residues
XX and is specifically recognised by autoimmune antibodies from patients
XX suffering from rheumatoid arthritis. The peptide comprises amino acids
XX involved in side chain interactions which is essential for the formation
XX of three-dimensional structure of the peptide. The peptide of the
XX invention is useful as a medicament to treat autoimmune diseases,
XX preferably rheumatoid arthritis. It is also useful for treating
XX autoimmune diseases by increasing the size of antigen-immune complexes to
XX improve clearance of the formed immune complexes and for the preparation
XX of a medicament for oral or nasal administration to treat autoimmune
XX diseases by inducing a state of systemic hyporesponsiveness or tolerance
XX to the peptide
XX
XX Sequence 14 AA;
XX
Query Match 41.9%; Score 44; DB 4; Length 14;
Best Local Similarity 76.9%; Pred. NO. 34;
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 6 GWGCDXSGRPGQ 18
DB 2 GHGCDXSGHRCGQ 14
XX
RESULT 7
AAE07382
ID AAE07382 standard; peptide; 16 AA.
XX
AC AAE07382;
XX
XX 10-APR-2001 (first entry)
XX
DE ErbB2 binding peptide amino acid sequence SEQ ID 33.
XX
XX Human; erbB2; HER2; cancer; nervous system disease; stroke; ischaemia;
XX metabolic disorder; nutritional deficiency; Alzheimer's disease;
XX Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
XX Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.
XX
OS Synthetic.
XX
XX WO200101748-A2.
XX
XX 11-JAN-2001.
XX
XX 30-JUN-2000; 2000WO-US018283.
XX
XX 02-JUL-1999; 99US-0142232P.
XX
XX (GETH ) GENENTECH INC.
XX
XX Dennis MS;
XX
XX WPI; 2001-123048/13.
XX
XX Non-naturally occurring peptide ligands which compete for binding human
XX ErbB2 gene products, useful for treating e.g. Alzheimer's disease,
XX multiple sclerosis and diabetic neuropathy.
XX
XX Disclosure; Fig 16; 116pp; English.
XX
XX This invention relates to non-naturally occurring peptide ligands which
XX bind to the human erbB2 gene product ErbB2 (also known as HER2). Peptides
XX represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
XX of the ErbB2 binding ligands of the invention. Sequences AAB76421 -
XX AAB76431 represent antibody Fc amino acid sequences used in the isolation
XX of the peptides of the invention. The peptides compete for binding ErbB2
XX with naturally occurring ligands, and may be used to treat disorders
XX characterized by over expression of HER2/ErbB2 such as cancers, diseases
XX of the nervous system, musculature and epithelia, e.g. nervous system
XX damage resulting from trauma, surgery, strokes, ischaemia, infection,
XX metabolic disorders, nutritional deficiency or toxic agents. In
XX particular the synthetic peptide ligands may be used to treat Alzheimer's
XX disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
XX chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
XX associated with diabetes
XX
XX Sequence 16 AA;
XX
Query Match 41.9%; Score 44; DB 4; Length 16;
Best Local Similarity 60.0%; Pred. NO. 38;
Matches 6; Conservative 1; Mismatches 3; Indels 0; Gaps 0;
QY 6 GWGCDXSGCR 15
DB 5 GWGCI GPCK 14
XX
RESULT 8
AAE07227
ID AAE07227 standard; peptide; 14 AA.
XX
AC AAE07227;
XX
XX 06-NOV-2001 (first entry)
XX
XX IGP1676 peptide for diagnosis and treatment of rheumatoid arthritis.
XX Synthetic peptide; cyclic; IGP1676; autoimmune antibody;
XX rheumatoid arthritis; therapy; autoimmune disease; anti-rheumatic;
XX
XX

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KW systemic hyporesponsiveness; immunosuppressive; antiarthritic.

XX Synthetic.

XX Key Location/Qualifiers
 XX Modified-site 1. 14
 FT /note= "Biotinylated residues"
 FT Disulfide-bond 9. 14
 FT Modified-site 11
 FT /note= "Citrulline"
 FT Modified-site 12
 FT /note= "Citrulline"

XX WO200146222-A2.

XX 28-JUN-2001.

XX 20-DEC-2000; 2000WO-EP013037.

XX 21-DEC-1999; 99EP-00870280.

XX 08-SEP-2000; 2000EP-00870195.

XX (INNO-) INNOGENETICS NV.

XX Union A, Moereels H, Meheus L;

XX WPI; 2001-496657/54.

XX New peptides, useful for diagnosing and treating rheumatoid arthritis,
 PT comprises citrulline residue between 2 cysteine residues and is
 PT specifically recognized by autoimmune antibodies from patients suffering
 PT from rheumatoid arthritis.

XX Claim 9; Page 42; 53pp; English.

XX The present sequence is a cyclic synthetic biotinylated peptide, IGP1676.
 CC The peptide comprises a citrulline residue between 2 cysteine residues
 CC and is specifically recognised by autoimmune antibodies from patients
 CC suffering from rheumatoid arthritis. The peptide comprises amino acids
 CC involved in side chain interactions which is essential for the formation
 CC of three-dimensional structure of the peptide. The peptide of the
 CC invention is useful as a medicament to treat autoimmune diseases,
 CC preferably rheumatoid arthritis. It is also useful for treating
 CC autoimmune diseases by increasing the size of antigen-immune complexes to
 CC improve clearance of the formed immune complexes and for the preparation
 CC of a medicament for oral or nasal administration to treat autoimmune
 CC diseases by inducing a state of systemic hyporesponsiveness or tolerance
 CC to the peptide

XX Sequence 14 AA;

Query Match 41.0%; Score 43; DB 4; Length 14;
 Best Local Similarity 66.7%; Pred. NO. 46;
 Matches 8; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 6 GWGCDXGCRPG 17

Db 2 GHPCXXGCRPG 13

RESULT 9

AAE07223

ID AAE07223 standard; peptide; 18 AA.

XX AAE07223;

XX 06-NOV-2001 (first entry)

XX IGP1648 peptide for diagnosis and treatment of rheumatoid arthritis.

XX Synthetic peptide; cyclic; IGP1648; autoimmune antibody;

XX rheumatoid arthritis; therapy; autoimmune disease; arthritic;

KW systemic hyporesponsiveness; immunosuppressive; antiarthritic.

XX Synthetic.

XX Key Location/Qualifiers
 XX Modified-site 1. 18
 FT /note= "Biotinylated residues"
 FT Disulfide-bond 9. 16
 FT Modified-site 11
 FT /note= "Citrulline"
 FT Modified-site 12
 FT /note= "Citrulline"

XX WO200146222-A2.

XX 28-JUN-2001.

XX 20-DEC-2000; 2000WO-EP013037.

XX 21-DEC-1999; 99EP-00870280.

XX 08-SEP-2000; 2000EP-00870195.

XX (INNO-) INNOGENETICS NV.

XX Union A, Moereels H, Meheus L;

XX WPI; 2001-496657/54.

XX New peptides, useful for diagnosing and treating rheumatoid arthritis,
 PT comprises citrulline residue between 2 cysteine residues and is
 PT specifically recognized by autoimmune antibodies from patients suffering
 PT from rheumatoid arthritis.

XX Claim 9; Page 42; 53pp; English.

XX The present sequence is a cyclic synthetic biotinylated peptide, IGP1648.
 CC The peptide comprises a citrulline residue between 2 cysteine residues
 CC and is specifically recognised by autoimmune antibodies from patients
 CC suffering from rheumatoid arthritis. The peptide comprises amino acids
 CC involved in side chain interactions which is essential for the formation
 CC of three-dimensional structure of the peptide. The peptide of the
 CC invention is useful as a medicament to treat autoimmune diseases,
 CC preferably rheumatoid arthritis. It is also useful for treating
 CC autoimmune diseases by increasing the size of antigen-immune complexes to
 CC improve clearance of the formed immune complexes and for the preparation
 CC of a medicament for oral or nasal administration to treat autoimmune
 CC diseases by inducing a state of systemic hyporesponsiveness or tolerance
 CC to the peptide

XX Sequence 18 AA;

Query Match 41.0%; Score 43; DB 4; Length 18;
 Best Local Similarity 61.1%; Pred. No. 58;
 Matches 11; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 1 QDTIVGWGCDXGCRPG 18

Db 1 QDTIHGHPGCSXXHRCGQ 18

RESULT 10

AAB76392

ID AAB76392 standard; peptide; 16 AA.

XX AAB76392;

XX 10-APR-2001 (first entry)

XX ErbB2 binding peptide amino acid sequence SEQ ID 43.

XX Human; erbB2; HER2; cancer; nervous system disease; stroke; ischaemia;
 KW metabolic disorder; nutritional deficiency; Alzheimer's disease;
 KW Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
 KW Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.

```

XX OS Synthetic.
XX PN WO200101748-A2.
XX PD 11-JAN-2001.
XX PF 30-JUN-2000; 2000WO-US018283.
XX PR 02-JUL-1999; 99US-0142232P.
XX PA (GETH ) GENENTECH INC.
XX PI Dennis MS;
XX DR WPI; 2001-123048/13.
XX PT Non-naturally occurring peptide ligands which compete for binding human
XX PT Erb2 gene products, useful for treating e.g. Alzheimer's disease,
XX PT multiple sclerosis and diabetic neuropathy.
XX PS Disclosure; Fig 16; 116pp; English.
XX CC This invention relates to non-naturally occurring peptide ligands which
XX CC bind to the human erbB2 gene product ErbB2 (also known as HER2). Peptides
XX CC represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
XX CC of the ErbB2 binding ligands of the invention. Sequences AAB76421 -
XX CC AAB76431 represent antibody Fc amino acid sequences used in the isolation
XX CC of the peptides of the invention. The peptides compete for binding ErbB2
XX CC with naturally occurring ligands, and may be used to treat disorders
XX CC characterized by over expression of HER2/ErbB2 such as cancers, diseases
XX CC of the nervous system, musculature and epithelia, e.g. nervous system
XX CC damage resulting from trauma, surgery, strokes, ischaemia, infection,
XX CC metabolic disorders, nutritional deficiency or toxic agents. In
XX CC particular the synthetic peptide ligands may be used to treat Alzheimer's
XX CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
XX CC chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
XX CC associated with diabetes
XX SQ Sequence 16 AA;
      Query Match 40.0%; Score 42; DB 4; Length 16;
      Best Local Similarity 66.7%; Pred. No. 71;
      Matches 6; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 6 GWGCDXGXC 14
DB |||||
5 GWGCIQPGC 13

RESULT 11
AAB76387
ID AAB76387 standard; peptide; 16 AA.
XX AC AAB76387;
XX 10-APR-2001 (first entry)
XX DE ErbB2 binding peptide amino acid sequence SEQ ID 38.
XX KW Human; erbB2; HER2; cancer; nervous system disease; stroke; ischaemia;
XX KW metabolic disorder; nutritional deficiency; Alzheimer's disease;
XX KW Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
XX KW Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.
XX OS Synthetic.
XX PN WO200101748-A2.
XX PD 11-JAN-2001.
XX PF 30-JUN-2000; 2000WO-US018283.
XX PR 02-JUL-1999; 99US-0142232P.
XX PA (GETH ) GENENTECH INC.
XX PI Dennis MS;
XX DR WPI; 2001-123048/13.
XX PT Non-naturally occurring peptide ligands which compete for binding human
XX PT Erb2 gene products, useful for treating e.g. Alzheimer's disease,
XX PT multiple sclerosis and diabetic neuropathy.
XX PS Disclosure; Fig 16; 116pp; English.
XX CC This invention relates to non-naturally occurring peptide ligands which
XX CC bind to the human erbB2 gene product ErbB2 (also known as HER2). Peptides
XX CC represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
XX CC of the ErbB2 binding ligands of the invention. Sequences AAB76421 -
XX CC AAB76431 represent antibody Fc amino acid sequences used in the isolation
XX CC of the peptides of the invention. The peptides compete for binding ErbB2
XX CC with naturally occurring ligands, and may be used to treat disorders
XX CC characterized by over expression of HER2/ErbB2 such as cancers, diseases
XX CC of the nervous system, musculature and epithelia, e.g. nervous system
XX CC damage resulting from trauma, surgery, strokes, ischaemia, infection,
XX CC metabolic disorders, nutritional deficiency or toxic agents. In
XX CC particular the synthetic peptide ligands may be used to treat Alzheimer's
XX CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
XX CC chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
XX CC associated with diabetes
XX SQ Sequence 16 AA;
      Query Match 40.0%; Score 42; DB 4; Length 16;
      Best Local Similarity 66.7%; Pred. No. 71;
      Matches 6; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 6 GWGCDXGXC 14
DB |||||
5 GWGCIQPGC 13

RESULT 12
AAB76369
ID AAB76369 standard; peptide; 17 AA.
XX AC AAB76369;
XX 10-APR-2001 (first entry)
XX DE ErbB2 binding peptide amino acid sequence SEQ ID 20.
XX KW Human; erbB2; HER2; cancer; nervous system disease; stroke; ischaemia;
XX KW metabolic disorder; nutritional deficiency; Alzheimer's disease;
XX KW Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
XX KW Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.
XX OS Synthetic.
XX PN WO200101748-A2.
XX PD 11-JAN-2001.
XX PF 30-JUN-2000; 2000WO-US018283.
XX PR 02-JUL-1999; 99US-0142232P.
XX PA (GETH ) GENENTECH INC.
XX PI Dennis MS;
XX DR WPI; 2001-123048/13.
XX PT Non-naturally occurring peptide ligands which compete for binding human

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PR 02-JUL-1999; 99US-0142232P.
XX (GETH ) GENENTECH INC.
XX PI Dennis MS;
XX DR WPI; 2001-123048/13.
XX PT Non-naturally occurring peptide ligands which compete for binding human
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XX PT multiple sclerosis and diabetic neuropathy.
XX PS Disclosure; Fig 16; 116pp; English.
XX CC This invention relates to non-naturally occurring peptide ligands which
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XX CC represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
XX CC of the ErbB2 binding ligands of the invention. Sequences AAB76421 -
XX CC AAB76431 represent antibody Fc amino acid sequences used in the isolation
XX CC of the peptides of the invention. The peptides compete for binding ErbB2
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XX CC characterized by over expression of HER2/ErbB2 such as cancers, diseases
XX CC of the nervous system, musculature and epithelia, e.g. nervous system
XX CC damage resulting from trauma, surgery, strokes, ischaemia, infection,
XX CC metabolic disorders, nutritional deficiency or toxic agents. In
XX CC particular the synthetic peptide ligands may be used to treat Alzheimer's
XX CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
XX CC chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
XX CC associated with diabetes
XX SQ Sequence 16 AA;
      Query Match 40.0%; Score 42; DB 4; Length 16;
      Best Local Similarity 66.7%; Pred. No. 71;
      Matches 6; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 6 GWGCDXGXC 14
DB |||||
5 GWGCIQPGC 13

RESULT 12
AAB76369
ID AAB76369 standard; peptide; 17 AA.
XX AC AAB76369;
XX 10-APR-2001 (first entry)
XX DE ErbB2 binding peptide amino acid sequence SEQ ID 20.
XX KW Human; erbB2; HER2; cancer; nervous system disease; stroke; ischaemia;
XX KW metabolic disorder; nutritional deficiency; Alzheimer's disease;
XX KW Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
XX KW Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.
XX OS Synthetic.
XX PN WO200101748-A2.
XX PD 11-JAN-2001.
XX PF 30-JUN-2000; 2000WO-US018283.
XX PR 02-JUL-1999; 99US-0142232P.
XX PA (GETH ) GENENTECH INC.
XX PI Dennis MS;
XX DR WPI; 2001-123048/13.
XX PT Non-naturally occurring peptide ligands which compete for binding human

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PT erB2 gene products, useful for treating e.g. Alzheimer's disease,
PT multiple sclerosis and diabetic neuropathy.
XX
XX Disclosure; Fig 16; 116pp; English.
XX This invention relates to non-naturally occurring peptide ligands which
bind to the human erB2 gene product ErB2 (also known as HER2). Peptides
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of the ErB2 binding ligands of the invention. Sequences AAB76421 -
CC AAB76431 represent antibody Fc amino acid sequences used in the isolation
of the peptides of the invention. The peptides compete for binding ErB2
with naturally occurring ligands, and may be used to treat disorders
characterized by over expression of HER2/ErB2 such as cancers, diseases
of the nervous system, musculature and epithelia, e.g. nervous system
damage resulting from trauma, surgery, strokes, ischaemia, infection,
CC metabolic disorders, nutritional deficiency or toxic agents. In
particular the synthetic peptide ligands may be used to treat Alzheimer's
disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
associated with diabetes
XX
SQ Sequence 17 AA;
Query Match 40.0%; Score 42; DB 4; Length 17;
Best Local Similarity 66.7%; Pred. No. 75;
Matches 6; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 6 GWGCDXSGC 14
|||||
DB 2 GWGIGPGC 10

RESULT 13
AAB76370
ID AAB76370 standard; peptide; 17 AA.
XX
AC AAB76370;
XX
DT 10-APR-2001 (first entry)
XX
DE ErB2 binding peptide amino acid sequence SEQ ID 21.
XX
XX Human; erB2; HER2; cancer; nervous system disease; stroke; ischaemia;
metabolic disorder; nutritional deficiency; Alzheimer's disease;
Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.
OS Synthetic.
XX
XX WO200101748-A2.
XX
XX 11-JAN-2001.
XX
XX 30-JUN-2000; 2000WO-US018283.
XX
XX 02-JUL-1999; 99US-0142232P.
XX
XX (GETH) GENENTECH INC.
XX
XX Dennis MS;
XX
XX WPI; 2001-123048/13.
XX
XX Non-naturally occurring peptide ligands which compete for binding human
erB2 gene products, useful for treating e.g. Alzheimer's disease,
multiple sclerosis and diabetic neuropathy.
XX
XX Disclosure; Fig 16; 116pp; English.
XX This invention relates to non-naturally occurring peptide ligands which
bind to the human erB2 gene product ErB2 (also known as HER2). Peptides
represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
of the ErB2 binding ligands of the invention. Sequences AAB76421 -

CC AAB76431 represent antibody Fc amino acid sequences used in the isolation
of the peptides of the invention. The peptides compete for binding ErB2
with naturally occurring ligands, and may be used to treat disorders
characterized by over expression of HER2/ErB2 such as cancers, diseases
of the nervous system, musculature and epithelia, e.g. nervous system
damage resulting from trauma, surgery, strokes, ischaemia, infection,
CC metabolic disorders, nutritional deficiency or toxic agents. In
particular the synthetic peptide ligands may be used to treat Alzheimer's
disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
associated with diabetes
XX
SQ Sequence 17 AA;
Query Match 40.0%; Score 42; DB 4; Length 17;
Best Local Similarity 66.7%; Pred. No. 75;
Matches 6; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 6 GWGCDXSGC 14
|||||
DB 2 GWGIGPGC 10

RESULT 14
AAE07221
ID AAE07221 standard; peptide; 18 AA.
XX
AC AAE07221;
XX
DT 06-NOV-2001 (first entry)
XX
DE IGP1646 peptide for diagnosis and treatment of rheumatoid arthritis.
XX
XX Synthetic peptide; cyclic; IGP1646; autoimmune antibody;
rheumatoid arthritis; therapy; autoimmune disease; anti-rheumatic;
XX systemic hyporesponsiveness; immunosuppressive; antiarthritic.
OS Synthetic.
XX
XX Key Location/Qualifiers
FT Modified-site 1.18 /note= "Biotinylated residues"
FT Disulfide-bond 9.16
FT Modified-site 12 /note= "Citrulline"
XX
XX WO200146222-A2.
XX
XX 28-JUN-2001.
XX
XX 20-DEC-2000; 2000WO-EP013037.
XX
XX 21-DEC-1999; 99EP-00870280.
XX
XX 08-SEP-2000; 2000EP-00870195.
XX
XX (INNO-) INNOGENETICS NV.
XX
XX Union A, Moereels H, Meheus L;
XX
XX WPI; 2001-496657/54.
XX
XX New peptides, useful for diagnosing and treating rheumatoid arthritis,
comprises citrulline residue between 2 cysteine residues and is
specifically recognized by autoimmune antibodies from patients suffering
from rheumatoid arthritis.
XX
XX Claim 9; Page 42; 53pp; English.
XX The present sequence is a cyclic synthetic biotinylated peptide, IGP1646.
CC The peptide comprises a citrulline residue between 2 cysteine residues
and is specifically recognised by autoimmune antibodies from patients
suffering from rheumatoid arthritis. The peptide comprises amino acids
involved in side chain interactions which is essential for the formation

CC of three-dimensional structure of the peptide. The peptide of the
CC invention is useful as a medicament to treat autoimmune diseases,
CC preferably rheumatoid arthritis. It is also useful for treating
CC autoimmune diseases by increasing the size of antigen-immune complexes to
CC improve clearance of the formed immune complexes and for the preparation
CC of a medicament for oral or nasal administration to treat autoimmune
CC diseases by inducing a state of systemic hyporesponsiveness or tolerance
CC to the peptide
XX
SQ Sequence 18 AA;

Query Match 40.0%; Score 42; DB 4; Length 18;
Best Local Similarity 64.7%; Pred. No. 78;
Matches 11; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 QDTIVGWCDSXGCRPG 17
| | | | | | | | | | | | | | | |
DB 1 QDTIHGHPCSSXGHRCG 17

RESULT 15
AAB76383
ID AAB76383 standard; peptide; 16 AA.
XX
AC AAB76383;

XX
DT 10-APR-2001 (first entry)
XX
DE ErbB2 binding peptide amino acid sequence SEQ ID 34.

XX Human; erbB2; HER2; cancer; nervous system disease; stroke; ischaemia;
KW metabolic disorder; nutritional deficiency; Alzheimer's disease;
KW Parkinson's disease; epilepsy; multiple sclerosis; Huntington's chorea;
KW Down's syndrome; nerve deafness; Meniere's disease; diabetic neuropathy.
XX
OS Synthetic.

XX WO200101748-A2.

XX PD 11-JAN-2001.

XX PF 30-JUN-2000; 2000WO-US018283.

XX PR 02-JUL-1999; 99US-0142232P.

XX PA (GETH) GENENTECH INC.

XX PI Dennis MS;

XX DR WPI; 2001-123048/13.

XX PT Non-naturally occurring peptide ligands which compete for binding human
PT erbB2 gene products, useful for treating e.g. Alzheimer's disease,
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XX PS Disclosure; Fig 16; 116pp; English.

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CC bind to the human erbB2 gene product ErbB2 (also known as HER2). Peptides
CC represented in AAB76350 - AAB76420 and AAB76432 - AAB76509 are examples
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CC of the nervous system, musculature and epithelia, e.g. nervous system
CC damage resulting from trauma, surgery, strokes, ischaemia, infection,
CC metabolic disorders, nutritional deficiency or toxic agents. In
CC particular the synthetic peptide ligands may be used to treat Alzheimer's
CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
CC chorea, Down's syndrome, nerve deafness, Meniere's disease and neuropathy
CC associated with diabetes

XX

SQ Sequence 16 AA;

Query Match 39.0%; Score 41; DB 4; Length 16;
Best Local Similarity 66.7%; Pred. No. 96;
Matches 6; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 7 WGCDSXGCR 15
| | | | | | | |
DB 6 WGCIGFGCR 14

Search completed: March 13, 2004, 07:28:55
Job time : 54 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 13, 2004, 07:26:14 ; Search time 20 Seconds
(without alignments)
86.572 Million cell updates/sec

Title: US-09-747-029b-17

Perfect score: 105

Sequence: 1 QDTIVGWGCDXGCRPGQ 18

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 3214

Minimum DB seq length: 0

Maximum DB seq length: 18

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: Pir1.*

2: Pir2.*

3: Pir3.*

4: Pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	29	27.6	15	D48394	major fat-globule
2	27	25.7	17	A61211	anantin - Streptom
3	26	24.8	10	RHEGG	gonadoliberin - sh
4	26	24.8	10	RHSHG	gonadoliberin - sh
5	25	23.8	15	I78838	flt3 ligand isoform
6	24	22.9	9	AKLQIM	locustamycininhibiti
7	24	22.9	11	D45900	complement C3b rec
8	24	22.9	15	S39012	proteinase - Therm
9	24	22.9	17	A44560	terephthalate 1,2-
10	23	21.9	10	A60421	hyperrethaloemic
11	23	21.9	10	S08997	hyperrethaloemic
12	23	21.9	10	S08998	hyperrethaloemic
13	23	21.9	10	A26381	hyperrethaloemic
14	23	21.9	10	A31571	hyperrethaloemic
15	23	21.9	10	S33995	hyperrethaloemic
16	23	21.9	11	S33300	probable substance
17	23	21.9	14	PH0755	T-cell receptor be
18	23	21.9	16	I78533	gene agouti protein
19	22	21.0	8	I57018	gene cfr protein
20	22	21.0	9	S39437	D-amino-acid oxida
21	22	21.0	10	RHAQ1	gonadoliberin I -
22	22	21.0	10	A13687	caerulein-like pep
23	22	21.0	11	S60354	retinal oxidase -
24	22	21.0	12	A49033	T-cell receptor de
25	22	21.0	13	PT0305	Ig heavy chain CRD
26	22	21.0	15	B56046	urinary tract ston
27	22	21.0	16	A49761	locustapyrokinin -
28	22	21.0	17	A61334	trypsin (BC 3.4.21
29	21.5	20.5	16	I79565	hypothetical TCU3/

30 21 20.0 9 2 A57444 neuropeptide Grb-A
31 21 20.0 11 2 B49164 chromogranin-B - r
32 21 20.0 12 2 I40663 bma protein - Clos
33 21 20.0 14 2 B61309 lutropin beta chain
34 21 20.0 15 2 S08301 epidermal growth f
35 21 20.0 16 2 D36912 hypothetical prote
36 21 20.0 17 2 PH1357 Ig heavy chain DJ
37 21 20.0 18 2 S21669 iH-4-oxoquinoline
38 21 20.0 18 2 B49048 T-cell receptor be
39 20 19.0 8 2 A31570 angiotensin-conver
40 20 19.0 8 2 JS0316 leucokinin VI - Ma
41 20 19.0 10 2 A61337 caerulein - frog
42 20 19.0 11 2 PT0273 Ig heavy chain CRD
43 20 19.0 11 2 PH0940 T-cell receptor be
44 20 19.0 12 2 B49033 T-cell receptor de
45 20 19.0 13 1 XAVI9B angiotensin-conver

ALIGNMENTS

RESULT 1

D48394

major fat-globule membrane protein GP 55 - guinea pig (fragment)

C;Species: Cavia porcellus (guinea pig)

C;Date: 19-Nov-1993 #sequence_revision 18-Nov-1994 #text_change 31-Oct-1997

C;Accession: D48394

R;Mather, I.H.; Banghart, L.R.; Lane, W.S.

Biochem. Mol. Biol. Int. 29, 545-554, 1993

A;Title: The major fat-globule membrane proteins, bovine components 15/16 and guinea-pi
II-like sequences.

A;Reference number: A48394; MUID:93250576; PMID:8485470

A;Accession: D48394

A;Status: preliminary

A;Molecule type: protein

A;Residues: 1-15 <MAT>

A;Experimental source: milk

A;Note: sequence extracted from NCBI backbone (NCBIP:131448)

C;Superfamily: milk fat globule protein; discoidin I amino-terminal homology; EGF homol

Query Match 27.6%; Score 29; DB 2; Length 15;

Best Local Similarity 57.1%; Pred. No. 5e+02;

Matches 4; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Qy 8 GCDXGC 14

Db 5 GCEIUGC 11

RESULT 2

A61211

anantin - Streptomyces coeruleus

C;Species: Streptomyces coeruleus

C;Date: 03-May-1994 #sequence_revision 05-Apr-1995 #text_change 07-May-1999

C;Accession: A61211

R;Wyss, D.F.; Lahm, H.W.; Manneberg, M.; Labhardt, A.M.

J. Antibiot. 44, 172-180, 1991

A;Title: Anantin -- a peptide antagonist of the atrial natriuretic factor (ANF). II. De

A;Reference number: A61211; MUID:91185186; PMID:1826288

A;Accession: A61211

A;Molecule type: protein

A;Residues: 1-17 <WYS>

A;Note: the isopeptide linked residue 8 is shown as Asn rather than Asp

F;1-8/Cross-link: isopeptide amino end (Gly-Asn) #status experimental

Query Match 25.7%; Score 27; DB 2; Length 17;

Best Local Similarity 44.4%; Pred. No. 1.1e+03;

Matches 4; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Qy 5 VWGCDXGC 13

Db 3 IGWGNIFG 11

RESULT 3

RHSGG
gonadoliberin - pig
C:Species: Sus scrofa domestica (domestic pig)
C>Date: 13-Jul-1981 #sequence_revision 13-Jul-1981 #text_change 18-Mar-1997
C/Accession: A01411
R:Baba, Y.; Matsuo, H.; Schally, A.V.
Biochem. Biophys. Res. Commun. 44, 459-463, 1971
A>Title: Structure of the porcine LH- and FSH-releasing hormone. II. Confirmation of the
A/Reference number: A90172; MUID:72114303; PMID:4946067
A/Accession: A01411
A/Molecule type: protein
A/Residues: 1-10 <BAB>
R:Matsuo, H.; Arimura, A.; Nair, R.M.G.; Schally, A.V.
Biochem. Biophys. Res. Commun. 45, 822-827, 1971
A>Title: Synthesis of the porcine LH- and FSH-releasing hormone by the solid-phase method
A/Reference number: A90176; MUID:72065376; PMID:4942726
A/Contents: annotation; synthesis
A/Note: the synthetic and natural hormones have the same physicochemical and biological
R:Baba, Y.; Arimura, A.; Schally, A.V.
Biochem. Biophys. Res. Commun. 45, 483-487, 1971
A>Title: On the tryptophan residue in porcine LH and FSH-releasing hormone.
A/Reference number: A90175; MUID:72117544; PMID:4946275
A/Contents: annotation
A/Note: Trp-3 appears to be essential for biological activity
C/Comment: This hypothalamic hormone stimulates the secretion of both luteinizing and follicle
C/Superfamily: gonadoliberin
C/Keywords: amidated carboxyl end; hormone; hypothalamus; pyroglutamic acid
F1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental
F10/Modified site: amidated carboxyl end (Gly) #status experimental

Query Match 24.8%; Score 26; DB 1; Length 10;
Best Local Similarity 71.4%; Pred. No. 9.7e+02; Mismatches 2; Indels 0; Gaps 0;
Matches 5; Conservative 0;

Qy 11 SXGCRPG 17
| | | | |
Db 4 SYGLRPG 10

RESULT 4

RHSHG
gonadoliberin - sheep
C:Species: Ovis orientalis aries, Ovis ammon aries (domestic sheep)
C>Date: 31-Dec-1991 #sequence_revision 31-Dec-1991 #text_change 16-Mar-1997
C/Accession: A93780; A01411
R:Burgus, R.; Butcher, M.; Amoss, M.; Ling, N.; Monahan, M.; Rivier, J.; Fellows, R.; Bl
Proc. Natl. Acad. Sci. U.S.A. 69, 278-282, 1972
A>Title: Primary structure of the ovine hypothalamic luteinizing hormone-releasing factor
A/Reference number: A93780; MUID:72094314; PMID:4550508
A/Accession: A93780
A/Molecule type: protein
A/Residues: 1-10 <SHR>
A/Note: the natural and synthetic hormones have the same biological activity
C/Comment: This hypothalamic hormone stimulates the secretion of both luteinizing and follicle
C/Superfamily: gonadoliberin
C/Keywords: amidated carboxyl end; hormone; hypothalamus; pyroglutamic acid
F1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental
F10/Modified site: amidated carboxyl end (Gly) #status experimental

Query Match 24.8%; Score 26; DB 1; Length 10;
Best Local Similarity 71.4%; Pred. No. 9.7e+02; Mismatches 2; Indels 0; Gaps 0;
Matches 5; Conservative 0;

Qy 11 SXGCRPG 17
| | | | |
Db 4 SYGLRPG 10

RESULT 5

I78838
flt3 ligand isoform E6 - mouse (fragment)

C:Species: Mus musculus (house mouse)
C>Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 05-Nov-1999
C/Accession: I78838
R:Lyman, S.D.; James, L.; Escobar, S.; Downey, H.; de Vries, P.; Brasel, K.; Stocking,
Oncogene 10, 149-157, 1995
A>Title: Identification of soluble and membrane-bound isoforms of the murine flt3 ligand
A/Reference number: I58343; MUID:95124710; PMID:7824267
A/Accession: I78838
A/Status: preliminary; translated from GB/EMBL/DBDJB
A/Molecule type: mRNA
A/Residues: 1-15 <RES>
A/Cross-references: GB:I576461; MID:9913481; PIDN:AAB33070.1; PID:9913482

Query Match 23.8%; Score 25; DB 2; Length 15;
Best Local Similarity 44.4%; Pred. No. 1.9e+03; Mismatches 4; Indels 0; Gaps 0;
Matches 4; Conservative 1;

Qy 9 CDSXGCRPG 17
| | | | |
Db 2 CLEUQCQPG 10

RESULT 6

AKQIM
locustamyoinhibiting peptide - migratory locust
C:Species: Locusta migratoria (migratory locust)
C>Date: 31-Mar-1993 #sequence_revision 31-Mar-1993 #text_change 20-Mar-1998
C/Accession: A60065
R:Schoofs, L.; Holman, G.M.; Hayes, T.K.; Nachman, R.J.; De Loof, A.
Regul. Pept. 36, 111-119, 1991
A>Title: Isolation, identification and synthesis of locustamyoinhibiting peptide (LOM-M
A/Reference number: A60065; MUID:92179466; PMID:1796179
A/Accession: A60065
A/Molecule type: protein
A/Residues: 1-9 <SCH>
C/Comment: This peptide hormone suppresses spontaneous contractions of the hindgut and
C/Superfamily: locustamyoinhibiting peptide
C/Keywords: amidated carboxyl end; hormone
F9/Modified site: amidated carboxyl end (Trp) #status experimental

Query Match 22.9%; Score 24; DB 1; Length 9;
Best Local Similarity 57.1%; Pred. No. 2.8e+05; Mismatches 3; Indels 0; Gaps 0;
Matches 4; Conservative 0;

Qy 1 QDTIVGV 7
| | | | |
Db 3 QDLNAGW 9

RESULT 7

D45900
complement C3b receptor type 2 - mouse (clone 12) (fragment)
C:Species: Mus musculus (house mouse)
C>Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change 07-May-1999
C/Accession: D45900
R:Kurtz, C.B.; O'Toole, E.; Christensen, S.M.; Weis, J.H.
J. Immunol. 144, 3581-3591, 1990
A>Title: The murine complement receptor gene family. IV. Alternative splicing of Cr2 gene
A/Reference number: A45900; MUID:90329754; PMID:2139460
A/Accession: D45900
A/Status: preliminary; nucleic acid sequence not shown; not compared with conceptual tr
A/Molecule type: mRNA
A/Residues: 1-11 <KUR>

Query Match 22.9%; Score 24; DB 2; Length 11;
Best Local Similarity 37.5%; Pred. No. 2e+03; Mismatches 3; Conservative 1; Indels 0; Gaps 0;
Matches 3; Conservative 1;

Qy 9 CDSXGCRPG 16
| | | | |
Db 2 CBEISCDP 9

RESULT 8
S39012
Proteinase - Thermus sp.
C:Species: Thermus sp.
C>Date: 18-Feb-1994 #sequence_revision 19-Apr-1996 #text_change 07-May-1999
C:Accession: S39012
R:Freeman, S.A.; Peek, K.; Prescott, M.; Daniel, R.
Biochem. J. 295, 463-469, 1993
A:Title: Characterization of a chelator-resistant proteinase from Thermus strain Rt4A2.
A:Reference number: S39012; MUID:94058984; PMID:8240244
A:Accession: S39012
A>Status: preliminary
A:Molecule type: protein
A:Residues: 1-15 <PRE>
A>Note: 13-Ala was also found

Query Match 22.9%; Score 24; DB 2; Length 15;
Best Local Similarity 66.7%; Pred. No. 2.7e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 VQWQCD 10
DB 6 VTWGLD 11

RESULT 9
A44560
terephthalate 1,2-dioxygenase (EC 1.-.-.-) oxygenase component alpha chain - Comamonas t
C:Species: Comamonas testosteroni
C>Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 03-Feb-1994
C:Accession: A44560
R:Schlaefli, H.
submitted to the Protein Sequence Database, December 1993
A:Description: Terephthalate 1,2-dioxygenase System from Comamonas testosteroni T-2: pur
A:Reference number: A44560
A:Accession: A44560
A:Molecule type: protein
A:Residues: 1-17 <SCH>
A>Note: it is uncertain whether the residue at position 9 is His or Arg
C:Keywords: oxidoreductase

Query Match 22.9%; Score 24; DB 2; Length 17;
Best Local Similarity 42.9%; Pred. No. 3e+03;
Matches 3; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 QDTIVGW 7
DB 2 QESIIQW 8

RESULT 10
A60421
hypertrehalosemic hormone - German cockroach
N:Alternate names: Bld-HrTH
C:Species: Blattella germanica (German cockroach)
C>Date: 03-Feb-1993 #sequence_revision 03-Feb-1993 #text_change 31-Oct-1997
C:Accession: A60421; S09137
R:Veenstra, J.A.; Camps, F.
Neuropeptides 15, 107-109, 1990
A:Title: Structure of the hypertrehalosemic neuropeptide of the German cockroach, Blatte
A:Reference number: A60421; MUID:91179584; PMID:2080017
A:Accession: A60421
A:Molecule type: protein
A:Residues: 1-10 <VEE>
R:Gaede, G.; Rinehart, K.L.
Biochem. Hoppe-Seyler 371, 345-354, 1990
A:Title: Primary structures of hypertrehalosemic neuropeptides isolated from the corpora
entalis and of the stick insect Extatosoma tiaratum assigned by tandem fast atom bombard
A:Reference number: S08995; MUID:90253659; PMID:2340112
A:Accession: S09137
A:Molecule type: protein
A:Residues: 1-10 <GAE>
C:Superfamily: adipokinetic hormone

C:Keywords: amidated carboxyl end; hormone; neuropeptide; pyroglutamic acid
F:1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental
F:10/Modified site: amidated carboxyl end (Thr) #status experimental

Query Match 21.9%; Score 23; DB 2; Length 10;
Best Local Similarity 100.0%; Pred. No. 2.6e+03;
Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GWG 8
DB 7 GWG 9

RESULT 11
S08997
hypertrehalosemic neuropeptide Bld-HrTH - cockroach (Gromphadorina portentosa)
C:Species: Gromphadorina portentosa
C>Date: 30-Jun-1992 #sequence_revision 14-Sep-1994 #text_change 24-Oct-1997
C:Accession: S08997
R:Gaede, G.; Rinehart, K.L.
Biochem. Hoppe-Seyler 371, 345-354, 1990
A:Title: Primary structures of hypertrehalosemic neuropeptides isolated from the corpora
entalis and of the stick insect Extatosoma tiaratum assigned by tandem fast atom bombard
A:Reference number: S08995; MUID:90253659; PMID:2340112
A:Accession: S08997
A:Molecule type: protein
A:Residues: 1-10 <GAE>
C:Superfamily: adipokinetic hormone
C:Keywords: amidated carboxyl end; hormone; neuropeptide; pyroglutamic acid
F:1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental
F:10/Modified site: amidated carboxyl end (Thr) #status experimental

Query Match 21.9%; Score 23; DB 2; Length 10;
Best Local Similarity 100.0%; Pred. No. 2.6e+03;
Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GWG 8
DB 7 GWG 9

RESULT 12
S08998
hypertrehalosemic neuropeptide Bld-HrTH - Madeira cockroach
C:Species: Leucophaea maderae (Madeira cockroach)
C>Date: 30-Jun-1992 #sequence_revision 14-Sep-1994 #text_change 24-Oct-1997
C:Accession: S08998
R:Gaede, G.; Rinehart, K.L.
Biochem. Hoppe-Seyler 371, 345-354, 1990
A:Title: Primary structures of hypertrehalosemic neuropeptides isolated from the corpora
entalis and of the stick insect Extatosoma tiaratum assigned by tandem fast atom bombard
A:Reference number: S08995; MUID:90253659; PMID:2340112
A:Accession: S08998
A:Molecule type: protein
A:Residues: 1-10 <GAE>
C:Superfamily: adipokinetic hormone
C:Keywords: amidated carboxyl end; hormone; neuropeptide; pyroglutamic acid
F:1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental
F:10/Modified site: amidated carboxyl end (Thr) #status experimental

Query Match 21.9%; Score 23; DB 2; Length 10;
Best Local Similarity 100.0%; Pred. No. 2.6e+03;
Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GWG 8
DB 7 GWG 9

RESULT 13
A26381
hypertrehalosemic hormone - gray cockroach
C:Species: Nauphoeta cinerea (gray cockroach)

C>Date: 31-Mar-1988 #sequence_revision 24-Oct-1997 #text_change 31-Oct-1997
 C/Accession: A26381
 R/Gade, G.; Rinehart Jr., K.L. 141, 774-781, 1986
 Biochem. Biophys. Res. Commun. 141, 774-781, 1986
 A/Title: Amino acid sequence of a hypotrehalosemic neuropeptide from the corpus cardia
 A/Reference number: A26381; MUID:87100208; PMID:3801028
 A/Accession: A26381
 A/Molecule type: protein
 A/Residues: 1-10 <GAD>
 A/Note: the amino-terminal residue forms pyrrolidone carboxylic acid; therefore, we have
 C/Superfamily: adipokinetic hormone
 C/Keywords: amidated carboxyl end; hormone; neuropeptide; pyroglutamic acid
 F1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental
 F10/Modified site: amidated carboxyl end (Thr) #status experimental

QY 6 GWG 8
 |||
 Db 7 GWG 9

Search completed: March 13, 2004, 07:26:51
 Job time : 22 secs

Query Match 21.9%; Score 23; DB 2; Length 10;
 Best Local Similarity 100.0%; Pred. No. 2.6e+03;
 Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GWG 8
 |||
 Db 7 GWG 9

RESULT 14

A31571
 hypotrehalosemic/adipokinetic hormone - bollworm
 N/Alternate names: Hez-HrTH
 C/Species: Heliothis zea (bollworm, corn earworm, tomato fruitworm)
 C/Date: 30-Jun-1989 #sequence_revision 23-Mar-1995 #text_change 31-Oct-1997
 C/Accession: A31571
 R/Jaffe, H.; Raina, A.K.; Riley, C.T.; Fraser, B.A.; Bird, T.G.; Tseng, C.M.; Zhang, Y.S
 Biochem. Biophys. Res. Commun. 155, 344-350, 1988
 A/Title: Isolation and primary structure of a neuropeptide hormone from Heliothis zea w
 A/Reference number: A31571; MUID:88326324; PMID:3415690
 A/Accession: A31571
 A/Molecule type: protein
 A/Residues: 1-10 <JAF>
 C/Superfamily: adipokinetic hormone
 C/Keywords: amidated carboxyl end; corpora cardiaca; hormone; neuropeptide; pyroglutamic
 F1/Modified site: pyrrolidone carboxylic acid (Gln) #status experimental
 F10/Modified site: amidated carboxyl end (Asn) #status experimental

Query Match 21.9%; Score 23; DB 2; Length 10;
 Best Local Similarity 100.0%; Pred. No. 2.6e+03;
 Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GWG 8
 |||
 Db 7 GWG 9

RESULT 15

B33995
 hypotrehalosemic hormone - black horse fly
 C/Species: Tabanus atratus (black horse fly)
 C/Date: 23-Mar-1990 #sequence_revision 23-Mar-1990 #text_change 31-Oct-1997
 C/Accession: B33995
 R/Jaffe, H.; Raina, A.K.; Riley, C.T.; Fraser, B.A.; Nachman, R.J.; Vogel, V.W.; Zhang,
 Proc. Natl. Acad. Sci. U.S.A. 86, 8161-8164, 1989
 A/Title: Primary structure of two neuropeptide hormones with adipokinetic and hypotrehal
 A/Reference number: A33995; MUID:90046758; PMID:2813385
 A/Accession: B33995
 A/Molecule type: protein
 A/Residues: 1-10 <JAF>
 C/Superfamily: adipokinetic hormone
 C/Keywords: amidated carboxyl end; corpora cardiaca; hormone; neuropeptide; pyroglutamic
 F1/Modified site: pyrrolidone carboxylic acid (Gln) #status predicted
 F10/Modified site: amidated carboxyl end (Tyr) #status predicted

Query Match 21.9%; Score 23; DB 2; Length 10;
 Best Local Similarity 100.0%; Pred. No. 2.6e+03;
 Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OM protein - protein search, using sw model.

Run on: March 13, 2004, 07:26:14 ; Search time 11 Seconds
(without alignments)

85.206 Million cell updates/sec

Title: US-09-747-029B-17

Perfect score: 105

Sequence: 1 QDTIVGWGCDXGCRFGQ 18

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 1002

Minimum DB seq length: 0

Maximum DB seq length: 18

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	32	30.5	17	1 CXMB_CONPE	P58926 conus penna
2	32	30.5	17	1 CXMB_CONPE	P58927 conus penna
3	25	23.8	17	1 NEF_FV1J3	P12480 human immun
4	24	22.9	9	1 LMIP_LOCM1	P31799 locusta mig
5	24	22.9	15	1 TAL_TREBR	P34070 tremella br
6	23	21.9	10	1 HTF_HELZE	P16353 heliothis z
7	23	21.9	10	1 HTF_NAUCI	P10939 nauphoeta c
8	23	21.9	10	1 HTF_TABAT	P14596 tabanus atr
9	23	21.9	11	1 TKNA_SCVCA	P41333 scyllorhinu
10	23	21.9	17	1 PA2_AUSSU	P59066 austrelaps
11	22	21.0	10	1 GON1_ALLMI	P37041 alligator m
12	22	21.0	11	1 CA31_LITCI	P82089 litoria cit
13	22	21.0	11	1 CA32_LITCI	P82090 litoria cit
14	22	21.0	12	1 HCYB_MEGRI	Q10584 megathura c
15	22	21.0	15	1 NXSO_PSETE	P59073 pseudonaja
16	22	21.0	16	1 LPK1_LOCM1	P20404 locusta mig
17	22	21.0	18	1 SODM_MYCHA	P80582 mycobacteri
18	21	20.0	12	1 CXST_CONTE	P58846 conus texti
19	21	20.0	13	1 CXL4_CONMR	P58810 conus marmo
20	21	20.0	14	1 KPPI_SELMI	P25933 selenastrum
21	21	20.0	15	1 UC19_MAIZE	P80625 zea mays (m
22	21	20.0	16	1 BAI1_EUBSP	P32371 eubacterium
23	20	19.0	8	1 ACI_THUAL	P18691 thunnus alb
24	20	19.0	8	1 LCK6_LEUNA	P19988 leucophaea
25	20	19.0	10	1 CA12_LITCI	P82086 litoria cit
26	20	19.0	10	1 CABR_LITXA	P82064 litoria xan
27	20	19.0	10	1 GON1_CLUPA	P81749 clupea pall
28	20	19.0	13	1 BPPI_EOTPA	P01020 bothrops ja
29	20	19.0	14	1 MAST_VESBA	P21654 vespa basal
30	20	19.0	15	1 KPP2_SELMI	P25934 selenastrum
31	20	19.0	16	1 LEC_DELEI	P83511 delonix reg
32	20	19.0	17	1 PATS_ANASP	O52748 anabaena sp
33	19	18.1	12	1 CXA1_CONIM	P50983 conus imper

34 19 18.1 15 1 CX1B_CONBE P58624 conus betul
35 19 18.1 16 1 TRYP_FELCA P81071 felis silve
36 19 18.1 17 1 GAST_MACMU P33714 macaca mula
37 19 18.1 17 1 GPX4_PINPS P81087 pinus pinas
38 18.5 17.6 15 1 GUN2_PINPS P81107 pinus pinas
39 18 17.1 8 1 CCKN_MACEU P30369 macropus eu
40 18 17.1 9 1 FAR5_CALVO P41860 calliphora
41 18 17.1 10 1 GON2_CHICK P37043 gallus gall
42 18 17.1 10 1 GON3_ONCKE P20367 oncorhynch
43 18 17.1 10 1 GONL_SQUAC P27429 squalus aca
44 18 17.1 12 1 RFI_CONSP P58805 conus spuri
45 18 17.1 14 1 IF2G_RAT P81795 rattus norv

ALIGNMENTS

RESULT 1
CXMA_CONPE STANDARD; PRT; 17 AA.
ID AC P58926;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Mu-conotoxin Pn1VA.
OS Conus pennaceus (feathered cone).
OC Eukaryota; Metazoa; Mollusca; Gastropoda; Orthogastropoda;
OC Apogastropoda; Caenogastropoda; Sorbeoconcha; Hypsogastropoda;
OC Neogastropoda; Conoidea; Conidae; Conus.
OX NCBI_TaxID=37335;
RN [1]
RP SEQUENCE, AND MASS SPECTROMETRY.
RX MEDLINE=95337083; PubMed=7612605;
RA Fainzilber M., Nakamura T., Gaathon A., Lodder J.C., Kitz K.S.,
RA Burlingame A.L., Zlotkin E.;
RT "A new cysteine framework in sodium channel blocking conotoxins.";
RL Biochemistry 34:8649-8656(1995).
CC -!- FUNCTION: Mu-conotoxins bind and block voltage-sensitive sodium
CC channel. Blocks reversibly sodium channels in molluscan neurons,
CC but has no effect on sodium currents in bovine chromaffin cells or
CC in rat brain synaptosomes. Induces paralysis in bivalve mollusks
CC (Mytilus). No effect are observed on fish (Gambusia) and fly
CC larvae (Sarcophaga). Pn1VB is approximately 6 times more potent
CC than Pn1VA in blockade of the sodium current in Lymnaea neurons.
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- TISSUE SPECIFICITY: Expressed by the venom duct.
CC -!- MASS SPECTROMETRY: MW=1789.5; METHOD=LSIMS.
CC -!- SIMILARITY: BELONGS TO THE M-SUPERFAMILY OF CONOTOXINS. MU-TYPE
CC FAMILY.
KW Toxin; Neurotoxin; Ionic channel inhibitor; Sodium channel inhibitor.
FT SITE 4 4
FT DISULFID 1 12 BY SIMILARITY.
FT DISULFID 2 15 BY SIMILARITY.
FT DISULFID 8 17 BY SIMILARITY.
SQ SEQUENCE 17 AA; 1797 MW; F9B721E0E96B9D82 CRC64;
Query Match 30.5%; Score 32; DB 1; Length 17;
Best Local Similarity 46.2%; Pred. No. 47;
Matches 6; Conservative 0; Mismatches 3; Indels 4; Gaps 1;
QY 6 GW----GDSXQC 14
DB 5 GWTCLLGCSPCC 17
RESULT 2
CXMB_CONPE STANDARD; PRT; 17 AA.
ID AC P58927;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Mu-conotoxin Pn1VB.

OS Conus pennaceus (Feathered cone).
OC Eukaryota; Metazoa; Mollusca; Gastropoda; Orthogastropoda;
OC Apogastropoda; Caenogastropoda; Sorbeoconcha; Hypogastropoda;
OC Neogastropoda; Conoidea; Conidae; Conus.
RN NCBI_TaxID=37335;
RX [1]
RP SEQUENCE, AND MASS SPECTROMETRY.
RX MEDLINE=95337083; PubMed=7612605;
RA Fainzilber M., Nakamura T., Gaathon A., Lodder J.C., Kitz K.S.,
RA Burlingame A.L., Zlotkin E.;
RT "A new cysteine framework in sodium channel blocking conotoxins";
RL Biochemistry 34:8649-8656(1995).
RN [2]
RP PHARMACOLOGICAL CHARACTERIZATION.
RX MEDLINE=95346025; PubMed=7620628;
RA Hasson A., Fainzilber M., Zlotkin E., Spira M.B.;
RT "Electrophysiological characterization of a novel conotoxin that
RT blocks molluscan sodium channels";
RL Eur. J. Neurosci. 7:815-818(1995).
CC -!- FUNCTION: Mu-conotoxins bind and block voltage-sensitive sodium
CC channel. Blocks reversibly sodium channels in molluscan neurons,
CC but has no effect on sodium currents in bovine chromaffin cells or
CC in rat brain synaptosomes. Induces paralysis in bivalve mollusks
CC (Mytilus). No effect are observed on fish (Gambusia) and fly
CC larvae (Sarcophaga). Is approximately 6 times more potent than
CC pNVA in blockade of the sodium current in *Lymnaea* neurons.
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- TISSUE SPECIFICITY: Expressed by the venom duct.
CC -!- MASS SPECTROMETRY: MW=1862.8; METHOD=LSIMS.
CC -!- SIMILARITY: BELONGS TO THE M-SUPERFAMILY OF CONOTOXINS. MU-TYPE
CC FAMILY.
KW Toxin; Neurotoxin; Ionic channel inhibitor; Sodium channel inhibitor.
FT SITE 4 4 IMPORTANT FOR BINDING AND ACTIVITY (BY
FT SIMILARITY).
FT DISULFID 1 12 BY SIMILARITY.
FT DISULFID 2 15 BY SIMILARITY.
FT DISULFID 8 17 BY SIMILARITY.
SQ SEQUENCE 17 AA; 1870 MW; E40021E0E96B9D82 CRC64;
Query Match 30.5%; Score 32; DB 1; Length 17;
Best Local Similarity 46.2%; Pred. No. 47;
Matches 6; Conservative 0; Mismatches 3; Indels 4; Gaps 1;
QY 6 GW----GCDXSGC 14
DB 5 GWTCLGCSGPCG 17
RESULT 3
NEF HV1J3
ID NEF HV1J3 STANDARD; PRT; 17 AA.
AC P12480;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Negative factor (F-protein) (27 kDa protein) (3'ORF) (Fragment).
GN NEF.
OS Human immunodeficiency virus type 1 (JH3 isolate) (HIV-1).
OC Viruses; Retroviral viruses; Retroviridae; Lentivirus.
CX NCBI_TaxID=11694;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=89352108; PubMed=2669897;
RA Komiya N., Hattori N., Inoue J., Sakuma S., Kurimura T., Yoshida M.;
RT "Nucleotide sequences of gag and env genes of a Japanese isolate of
RT HIV-1 and their expression in bacteria";
RL AIDS Res. Hum. Retroviruses 5:411-419(1989).
CC -!- FUNCTION: NEF has GTPase, GTP-binding and autophosphorylating
CC activities. It seems to down-regulate the CD4(T4) antigen.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC -----
CC EMBL; M21138; AAB03527.1; -.
DR HIV; M21138; NEFSJH3.
DR InterPro; IPR001558; HIV_Nef.
DR Pfam; PF00469; F-protein; 1.
KW AIDS; Myristate; GTP-binding; Lipoprotein.
FT LIPID 2 2 N-myristoyl glycine (in host) (By
FT similarity).
FT NON_TER 17 17
SQ SEQUENCE 17 AA; 1901 MW; 65E8B3F26FEF921E CRC64;
Query Match 23.8%; Score 25; DB 1; Length 17;
Best Local Similarity 60.0%; Pred. No. 5.8e+02;
Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 TIVGV 7
DB 9 SVVGV 13
RESULT 4
LMIP LOCMI STANDARD; PRT; 9 AA.
ID LMIP LOCMI
AC P31799;
DT 01-JUL-1993 (Rel. 26, Created)
DT 01-JUL-1993 (Rel. 26, Last sequence update)
DT 01-OCT-1993 (Rel. 27, Last annotation update)
DE Locustamyoinhibiting peptide (LOM-MIP).
OS Locusta migratoria (migratory locust).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Orthopteroidea; Orthoptera; Caelifera; Acridoidea;
OC Acridoidea; Acrididae; Oedipodinae; Locusta.
CX NCBI_TaxID=7004;
RN [1]
RP SEQUENCE.
RX MEDLINE=92179466; PubMed=1796179;
RA Schoofs L., Holman G.M., Hayes T.K., Nachman R.J., de Loof A.;
RT "Isolation, identification and synthesis of locustamyoinhibiting
RT peptide (LOM-MIP), a novel biologically active neuro-peptide from
RT Locusta migratoria";
RL Regul. Pept. 36:111-119(1991).
CC -!- FUNCTION: Suppresses spontaneous contractions of the hindgut and
CC oviduct.
CC -!- TISSUE SPECIFICITY: Neurons located in two ventral cell clusters
CC in the suboesophageal ganglion.
DR PIR; A60065; AKLQIM.
KW Amidation; Neuropeptide.
FT MOD_RES 9 9 AMIDATION.
SQ SEQUENCE 9 AA; 1060 MW; 387D7DD4472AB6C3 CRC64;
Query Match 22.9%; Score 24; DB 1; Length 9;
Best Local Similarity 57.1%; Pred. No. 1.4e+05;
Matches 4; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 QDTIVGV 7
DB 3 QDLNAGV 9
RESULT 5
TAL TREER
ID TAL TREER STANDARD; PRT; 15 AA.
AC P34070;
DT 01-FEB-1994 (Rel. 28, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Tremorgen A-1.
OS Tremella brasiliensis (Jelly fungus).
OC Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Heterobasidiomycetes;

OC Tremellomycetidae; Tremellales; Tremellaceae; Tremella.
 OX NCBI_TaxID=29896;
 RN [1]
 RP SEQUENCE.
 RA Ishibashi Y., Sakagami Y., Isegai A., Suzuki A.;
 RT "Structures of Tremogens A-9291-I and A-9291-VIII: peptidyl sex
 RL hormones of Tremella brasiliensis.";
 RT Biochemistry 23:1399-1404(1994).
 CC -!- FUNCTION: Tremogogen A-I is produced by the a mating-type cells
 CC and induces formation of conjugation tubes in a mating-type cells.
 KW Phoromone; Prenylation; Lipoprotein.
 FT LIPID 15 15 S-farnesyl cysteine.
 SQ SEQUENCE 15 AA; 1339 MW; 3AABA4FC2D605333 CRC64;
 Query Match 22.9%; Score 24; DB 1; Length 15;
 Best Local Similarity 57.1%; Pred. No. 7e+02;
 Matches 4; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 Oy 8 GCDXKGC 14
 | | | |
 Db 9 GASSGGC 15
 RESULT 6
 HTF_HELZE
 ID -HTF_HELZE STANDARD; PRT; 10 AA.
 AC P16353;
 DT 01-AUG-1990 (Rel. 15, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Hypertrehalosaemic hormone (HeZ-HRTH).
 OS Heliothis zea (Corn earworm) (Bollworm).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Lepidoptera; Glossata; Ditrystia; Noctuoidea;
 OC Noctuidae; Heliothinae; Helicoverpa.
 OX NCBI_TaxID=7113;
 RN [1]
 RP SEQUENCE.
 RC TISSUE=Corpora cardiaca;
 RX MEDLINE=89326324; PubMed=3415690;
 RA Jaffe H., Raina A.K., Riley C.T., Fraser B.A., Bird T.G.,
 RA Tseng C.M., Zhang Y.S., Hayes D.K.;
 RT "Isolation and primary structure of a neuropeptide hormone from
 RT Heliothis zea with hypertrehalosaemic and adipokinetic activities.";
 RL Biochem. Biophys. Res. Commun. 155:344-350(1988).
 CC -!- FUNCTION: Hypertrehalosaemic factors are neuropeptides that
 CC elevate the level of trehalose in the hemolymph (trehalose is the
 CC major carbohydrate in the hemolymph of insects).
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- SIMILARITY: Belongs to the AKH / HRTH / RPCH family.
 DR PIR; A31571; A31571.
 DR InterPro; IPR002047; AKH.
 DR PROSITE; PS00256; AKH; 1.
 KW Neuropeptide; Amidation; Pyrrolidone carboxylic acid.
 FT MOD_RES 1 1 PYRROLIDONE CARBOXYLIC ACID.
 FT MOD_RES 10 10 AMIDATION.
 SQ SEQUENCE 10 AA; 1096 MW; 8E70367865A5B9D1 CRC64;
 Query Match 21.9%; Score 23; DB 1; Length 10;
 Best Local Similarity 100.0%; Pred. No. 7e+02;
 Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 6 GWG 8
 | | | |
 Db 7 GWG 9
 RESULT 7
 HTF_NAUCI
 ID -HTF_NAUCI STANDARD; PRT; 10 AA.
 AC P10939;
 DT 01-JUL-1989 (Rel. 11, Created)
 DT 01-FEB-1994 (Rel. 26, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Hypertrehalosaemic hormone (HTH) (Hypertrehalosaemic neuropeptide).
 OS Nauphoeta cinerea (Cinereous cockroach) (Gray cockroach),
 OS Leucophaea maderae (Madeira cockroach),
 OS Blattella germanica (German cockroach), and
 OS Gromphadorina portentosa (Madagascan hissing cockroach).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Orthopteroidea; Dictyoptera; Blattaria; Blaberoidea;
 OC Blaberidae; Nauphoeta.
 OX NCBI_TaxID=6990, 6988, 6973, 36953;
 RN [1]
 RP SEQUENCE.
 RC SPECIES=N.cinerea; TISSUE=Corpora cardiaca;
 RX MEDLINE=87100208; PubMed=3801028;
 RA Gaede G., Rinehart K.L. Jr.;
 RT "Amino acid sequence of a hypertrehalosaemic neuropeptide from the
 RT corpus cardiaca of the cockroach, Nauphoeta cinerea.";
 RL Biochem. Biophys. Res. Commun. 141:774-781(1986).
 RN [2]
 RP SEQUENCE.
 RC SPECIES=L.maderae, G.portentosa, and B.germanica;
 RX MEDLINE=90253659; PubMed=2340112;
 RA Gaede G., Rinehart K.L. Jr.;
 RT "Primary structures of hypertrehalosaemic neuropeptides isolated from
 RT the corpora cardiaca of the cockroaches Leucophaea maderae,
 RT Gromphadorina portentosa, Blattella germanica and Blattella orientalis
 RT and of the stick insect Extatosoma tiaratum assigned by tandem fast
 RT atom bombardment mass spectrometry.";
 RL Biol. Chem. Hoppe-Seyler 371:345-354(1990).
 RN [3]
 RP SEQUENCE.
 RC SPECIES=B.germanica;
 RX MEDLINE=91179584; PubMed=2080017;
 RA Veenstra J.A., Camps F.;
 RT "Structure of the hypertrehalosaemic neuropeptide of the German
 RT cockroach, Blattella germanica.";
 RL Neuropeptides 15:107-109(1990).
 CC -!- FUNCTION: Hypertrehalosaemic factors are neuropeptides that
 CC elevate the level of trehalose in the hemolymph (trehalose is the
 CC major carbohydrate in the hemolymph of insects).
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- SIMILARITY: Belongs to the AKH / HRTH / RPCH family.
 DR PIR; A26381; A26381.
 DR PIR; A60421; A60421.
 DR PIR; S08997; S08997.
 DR PIR; S08998; S08998.
 DR InterPro; IPR002047; AKH.
 DR PROSITE; PS00256; AKH; 1.
 KW Neuropeptide; Amidation; Pyrrolidone carboxylic acid.
 FT MOD_RES 1 1 PYRROLIDONE CARBOXYLIC ACID.
 FT MOD_RES 10 10 AMIDATION.
 SQ SEQUENCE 10 AA; 1092 MW; 056236786775B9C4 CRC64;
 Query Match 21.9%; Score 23; DB 1; Length 10;
 Best Local Similarity 100.0%; Pred. No. 7e+02;
 Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 6 GWG 8
 | | | |
 Db 7 GWG 9
 RESULT 8
 HTF_TABAT
 ID -HTF_TABAT STANDARD; PRT; 10 AA.
 AC P14596;
 DT 01-JAN-1990 (Rel. 13, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Hypertrehalosaemic factor (HOTH) (Dipteran corpora cardiaca factor II)
 DE (DCC II).
 OS Tabanus atratus (Horse fly).
 OC Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;

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OC Neoptera; Endopterygota; Diptera; Brachycera; Tabanomorpha; Tabanidae;
OC Tabanus.
OX NCBI_TaxID=7207;
RN [1]
RP SEQUENCE.
RX TISSUE=Corpora cardiaca;
RM MEDLINE=90046758; PubMed=2813385;
RA Jaffe H., Raina A.K., Riley C.T., Fraser B.A., Nachman R.J.,
RA Vogel V.W., Zhang Y.-S., Hayes D.K.,
RT "Primary structure of two neuropeptide hormones with adipokinetic and
RT hypotrehalosemic activity isolated from the corpora cardiaca of horse
RT flies (Diptera).";
RL Proc. Natl. Acad. Sci. U.S.A. 86:8161-8164(1989).
CC -!- FUNCTION: Hyperrehalosemic factors are neuropeptides that
CC elevate the level of trehalose in the hemolymph (trehalose is the
CC major carbohydrate in the hemolymph of insects).
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- SIMILARITY: Belongs to the AKH / HRTH / RPCH family.
DR PIR; B33995; B33995.
DR InterPro; IPR002047; AKH.
DR PROSITE; PS00256; AKH, 1.
KW Neuropeptide; Amidation; Pyrrolidone carboxylic acid.
FT MOD_RES 1 1 PYRROLIDONE CARBOXYLIC ACID.
FT MOD_RES 10 10 AMIDATION.
SQ SEQUENCE 10 AA; 1169 MW; 916036786771A9D1 CRC64;

Query Match 21.9%; Score 23; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 7e+02;
Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GWG 8
Db 7 GWG 9

RESULT 9
TKNA SCYCA STANDARD; PRT; 11 AA.
AC P41333;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Substance P.
OS Scyliorhinus canicula (Spotted dogfish) (Spotted catshark).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Chondrichthyes;
OC Elasmobranchii; Galeomorphii; Galeoidea; Carchariniiformes;
OC Scyliorhinidae; Scyliorhinus.
OX NCBI_TaxID=7830;
RN [1]
RP SEQUENCE.
RX TISSUE=Brain;
RM MEDLINE=93292508; PubMed=7685693;
RA Waugh D., Wang Y., Hazon N., Balmert R.J., Conlon J.M.;
RT "Primary structures and biological activities of substance-P-related
RT peptides from the brain of the dogfish, Scyliorhinus canicula.";
RL Eur. J. Biochem. 214:469-474(1993).
CC -!- FUNCTION: Tachykinins are active peptides which excite neurons,
CC evoke behavioral responses, are potent vasodilators and
CC secretagogues, and contract (directly or indirectly) many smooth
CC muscles.
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- SIMILARITY: Belongs to the tachykinin family.
DR PIR; S33300; S33300.
DR InterPro; IPR002040; Tachy. Neurokinin.
DR PROSITE; PS00267; TACHYKININ, 1.
KW Tachykinin; Neuropeptide; Amidation; Neurotransmitter.
FT MOD_RES 11 11 AMIDATION.
SQ SEQUENCE 11 AA; 1278 MW; 214860DEC9D6D867 CRC64;

Query Match 21.9%; Score 23; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 7.7e+02;
Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 15 RFGQ 18
Db 3 RFGQ 6

RESULT 10
PA2 AUSSU STANDARD; PRT; 17 AA.
AC P59066;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Phospholipase A2 (EC 3.1.1.4) (Phosphatidylcholine 2-acylhydrolase)
DE (Fragment).
OS Austrelaps superbus (Australian copperhead).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Lepidosauria; Squamata; Scleroglossa; Serpentes; Colubroidea;
OC Elapidae; Acanthophilineae; Austrelaps.
OX NCBI_TaxID=29156;
RN [1]
RP SEQUENCE.
RX MEDLINE=93369790; PubMed=8362372;
RA Yuan Y., Jackson S.P., Mitchell C.A., Salem H.H.;
RT "Purification and characterisation of a snake venom phospholipase A2:
RT a potent inhibitor of platelet aggregation.";
RL Thromb. Res. 70:471-481(1993).
CC -!- FUNCTION: PA2 catalyzes the calcium-dependent hydrolysis of the 2-
CC acyl groups in 3-sn-phosphoglycerides. Inhibits collagen-, ADP-,
CC thrombin-, ionophore-, adrenaline-, ristocetin-, and arachidonic
CC acid-induced platelet aggregation. Inhibits serotonin release.
CC -!- CATALYTIC ACTIVITY: Phosphatidylcholine + H(2)O = 1-
CC acyl-glycerophosphocholine + a fatty acid anion.
CC -!- COFACTOR: Calcium (Probable).
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- TISSUE SPECIFICITY: Expressed by the venom gland.
CC -!- SIMILARITY: Belongs to the phospholipase A2 family. Group I
CC subfamily.
DR InterPro; IPR001211; PhospholipaseA2.
DR Pfam; PF00668; phoslip; 1.
KW Lipid degradation; Hydrolase; Toxin; Calcium.
FT NON_TER 17 17
SQ SEQUENCE 17 AA; 1846 MW; 03FB7DD7B7D7D1CB CRC64;

Query Match 21.9%; Score 23; DB 1; Length 17;
Best Local Similarity 50.0%; Pred. No. 1.2e+03;
Matches 4; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 8 GCDXGCR 15
Db 10 GCANHGR 17

RESULT 11
GONI ALLMI STANDARD; PRT; 10 AA.
AC P37041; P20407;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-FEB-1991 (Rel. 17, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Gonadoliberin I (Gonadotropin-releasing hormone I) (GnRH-I) (LH-RH I)
DE (luliberin I).
OS Alligator mississippiensis (American alligator).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Crocodylidae; Alligatorinae; Alligator.
OX NCBI_TaxID=8496;
RN [1]
RP SEQUENCE.
RX TISSUE=Brain;
RM MEDLINE=91352338; PubMed=1882082;
RA Lovejoy D.A., Fischer W.H., Parker D.B., McRory J.E., Park M.,
RA Lance V., Swanson P., Rivier J.E., Sherwood N.M.;
RT "Primary structure of two forms of gonadotropin-releasing hormone

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RT from brains of the American alligator (Alligator mississippiensis).";
 RL Regul. Pept. 33:105-116(1991)
 CC -!- FUNCTION: Stimulates the secretion of gonadotropins.
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- SIMILARITY: Belongs to the GnRH family.

DR PIR; A60066; RHAQ1.
 DR InterPro; IPR002012; GnRH.
 DR Pfam; PF00446; GnRH; 1.
 DR PROSITE; PS00473; GnRH; 1.
 KW Hormone; Amidation; Hypothalamus; Pyrrolidone carboxylic acid.
 FT MOD_RES 1 1
 FT MOD_RES 10 10 PYRROLIDONE CARBOXYLIC ACID.
 FT AMIDATION.
 SQ SEQUENCE 10 AA; 1172 MW; 284923D7286B45A3 CRC64;

Query Match 21.0%; Score 22; DB 1; Length 10;
 Best Local Similarity 57.1%; Pred. No. 1e+03;
 Matches 4; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 11 SXGCRPG 17
 ||||
 DB 4 SYGLQPG 10

RESULT 12

CA31_LITCI
 ID CA31_LITCI STANDARD; PRT; 11 AA.

AC P82089; (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Caerulein 3.1/3.1Y4.
 OS Litoria citropa (Australian blue mountains tree frog).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Amphibia; Batrachia; Anura; Neobatrachia; Hylidae;
 CC Pelodyadinae; Litoria.
 CC NCBI_TaxID=94770;
 RN [1]

SEQUENCE, AND MASS SPECTROMETRY.

RP TISSUE=Skin secretion;
 RX MEDLINE=20057701; PubMed=10589099;
 RA Wabnitz P.A., Bowie J.H., Tyler M.J.;
 RT "Caerulein-like peptides from the skin glands of the Australian blue mountains tree frog Litoria citropa. Part 1. Sequence determination using electrospray mass spectrometry."
 RL Rapid Commun. Mass Spectrom. 13:2498-2502(1999).
 CC -!- FUNCTION: Hypotensive neuropeptide (Probable).
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- TISSUE SPECIFICITY: Skin dorsal glands.
 CC -!- PTM: Isoform 3.1Y4 differs from isoform 3.1 in not being sulfated.

CC -!- MASS SPECTROMETRY: MW=1407; METHOD=Electrospray.
 CC -!- SIMILARITY: Belongs to the gastrin/cholecystokinin family.
 DR InterPro; IPR001651; Gastrin.
 DR PROSITE; PS00259; GASTRIN; FALSE_NEG.
 KW Amphibian defense peptide; Hypotensive agent; Amidation; Sulfation;
 FT MOD_RES 1 1
 FT MOD_RES 4 4 PYRROLIDONE CARBOXYLIC ACID.
 FT MOD_RES 11 11 SULFATION.
 SQ SEQUENCE 11 AA; 1347 MW; 10DAB7D67861A86B CRC64;

Query Match 21.0%; Score 22; DB 1; Length 11;
 Best Local Similarity 57.1%; Pred. No. 1.1e+03;
 Matches 4; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 QDTIVGW 7
 ||||
 DB 2 QDYGTGW 8

RESULT 13

CA32_LITCI
 ID CA32_LITCI STANDARD; PRT; 11 AA.

AC P82090;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 10-OCT-2001 (Rel. 40, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Caerulein 3.2/3.2Y4.
 OS Litoria citropa (Australian blue mountains tree frog).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Amphibia; Batrachia; Anura; Neobatrachia; Hylidae;
 CC Pelodyadinae; Litoria.
 CC NCBI_TaxID=94770;
 RN [1]

SEQUENCE, AND MASS SPECTROMETRY.

RP TISSUE=Skin secretion;
 RX MEDLINE=20057701; PubMed=10589099;
 RA Wabnitz P.A., Bowie J.H., Tyler M.J.;
 RT "Caerulein-like peptides from the skin glands of the Australian blue mountains tree frog Litoria citropa. Part 1. Sequence determination using electrospray mass spectrometry."
 RL Rapid Commun. Mass Spectrom. 13:2498-2502(1999).
 CC -!- FUNCTION: Hypotensive neuropeptide (Probable).
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- TISSUE SPECIFICITY: Skin dorsal glands.
 CC -!- PTM: Isoform 3.2Y4 differs from isoform 3.2 in not being sulfated.

CC -!- MASS SPECTROMETRY: MW=1423; METHOD=Electrospray.
 CC -!- SIMILARITY: Belongs to the gastrin/cholecystokinin family.
 DR InterPro; IPR001651; Gastrin.
 DR PROSITE; PS00259; GASTRIN; FALSE_NEG.
 KW Amphibian defense peptide; Hypotensive agent; Amidation; Sulfation;
 KW Pyrrolidone carboxylic acid. PYRROLIDONE CARBOXYLIC ACID.
 FT MOD_RES 1 1
 FT MOD_RES 4 4 SULFATION.
 FT MOD_RES 11 11 AMIDATION.
 SQ SEQUENCE 11 AA; 1363 MW; 10DAB8867861A86B CRC64;

Query Match 21.0%; Score 22; DB 1; Length 11;
 Best Local Similarity 57.1%; Pred. No. 1.1e+03;
 Matches 4; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 QDTIVGW 7
 ||||
 DB 2 QDYGTGW 8

RESULT 14

HCYB_MEGCR
 ID HCYB_MEGCR STANDARD; PRT; 12 AA.

AC Q10584;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hemocyanin B chain (KLH-B) (Fragment).
 OS Megathura crenulata (Giant keyhole limpet).
 CC Eukaryota; Metazoa; Mollusca; Gastropoda; Orthogastropoda;
 CC Vetiagastropoda; Fissurelloidea; Fissurellidae; Megathura.
 CC NCBI_TaxID=55429;
 RN [1]

SEQUENCE

RP MEDLINE=96208935; PubMed=8929804;
 RA Swerdlow R.D., Ebert R.F., Lee P., Bonaventura C., Miller K.I.;
 RT "Keyhole limpet hemocyanin: structural and functional characterization of two different subunits and multimers."
 RL Comp. Biochem. Physiol. 113B:537-548(1996).
 CC -!- FUNCTION: Hemocyanins are copper-containing oxygen carriers occurring freely dissolved in the hemolymph of many mollusks and arthropods.

CC -!- SUBUNIT: Dodecamers and extended multimers.
 CC -!- SUBCELLULAR LOCATION: Extracellular.
 CC -!- TISSUE SPECIFICITY: Hemolymph.

CC -!- BIOTECHNOLOGY: Potent immunogen used classically as a carrier protein for haptens and more recently in human vaccines and for immunotherapy of bladder cancer.

CC -!- SIMILARITY: Belongs to the tyrosinase family. Hemocyanin

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CC      subfamily.
DR      InterPro; IPR000896; Hemocyanin.
DR      PROSITE; PS00209; HEMOCYANIN_1; PARTIAL.
DR      PROSITE; PS00210; HEMOCYANIN_2; PARTIAL.
KW      Oxygen transport; Transport; Copper; Glycoprotein;
KW      Hemolymph.
FT      NON_TER      12      12
SQ      SEQUENCE      12 AA; 1345 MW; CEFBEAA44A32412 CRC64;

Query Match      21.0%; Score 22; DB 1; Length 12;
Best Local Similarity 50.0%; Pred. NO. 1.2e+03;
Matches 5; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY      2 DTIVGWGCDGDS 11
      |||
      |||
      |||
DB      2 DIVVRKNVDS 11

RESULT 15
NXSO_PSET6
ID      NXSO_PSET6      STANDARD;      PRT;      15 AA.
AC      P59073;
DT      28-FEB-2003 (Rel. 41, Created)
DT      28-FEB-2003 (Rel. 41, Last sequence update)
DT      28-FEB-2003 (Rel. 41, Last annotation update)
DE      Short neurotoxin N2 (Alpha neurotoxin) [fragment].
OS      Pseudonaja textilis (Eastern brown snake).
OC      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC      Lepidosauria; Squamata; Chordata; Serpentes; Colubroides;
OC      Elapidae; Acanthophiinae; Pseudonaja.
OX      NCBI_TaxID=8673;
RN      [1]
RP      SEQUENCE, AND MASS SPECTROMETRY.
RC      TISSUE=Venom;
RX      MEDLINE=9449602; PubMed=10518793;
RA      Gong N.L., Armugam A., Jayaseelan K.;
RT      "Postsynaptic short-chain neurotoxins from Pseudonaja textilis: cDNA
RT      cloning, expression and protein characterization.";
RL      Eur. J. Biochem. 265:982-989(1999).
CC      -!- FUNCTION: Lethal neurotoxin, binds and inhibits nicotinic
CC      acetylcholine receptors (nAChR).
CC      -!- SUBCELLULAR LOCATION: Secreted.
CC      -!- TISSUE SPECIFICITY: Expressed by the venom gland.
CC      -!- MASS SPECTROMETRY: MW=6345; METHOD=Electrospray.
CC      -!- MISCELLANEOUS: LD(50) is 0.80 mg/kg by intravenous injection.
CC      -!- SIMILARITY: Belongs to the snake toxin family.
DR      InterPro; IPR003571; Snake toxin.
DR      PROSITE; PS00272; SNAKE_TOXIN; PARTIAL.
KW      Toxin; Neurotoxin; Postsynaptic neurotoxin;
KW      Acetylcholine receptor inhibitor; Multigene family.
FT      UNSURE      3      3
FT      UNSURE      13      13
FT      NON_TER      15      15
SQ      SEQUENCE      15 AA; 1727 MW; E149FD4BFD1EF0DD CRC64;

Query Match      21.0%; Score 22; DB 1; Length 15;
Best Local Similarity 33.3%; Pred. NO. 1.5e+03;
Matches 5; Conservative 2; Mismatches 0; Indels 8; Gaps 1;

QY      2 DTIVGWGCDGSGCRP 16
      |||
      |||
      |||
DB      9 DIVVRKNVDS 11

Search completed: March 13, 2004, 07:27:15
Job time : 13 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 13, 2004, 07:26:14 ; Search time 39 Seconds
(without alignments)
145.624 Million cell updates/sec

Title: US-09-747-029B-17

Perfect score: 105

Sequence: 1 QDTIVGWGCDXGCRPCQ 18

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 5675

Minimum DB seq length: 0

Maximum DB seq length: 18

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL 25:
1: sp_archea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phase:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_virus:*
16: sp_bacteriap:*
17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	34	32.4	15	3 Q14379	O14379 schizosacch
2	30	28.6	18	2 Q09053	Q09053 methylenona
3	27	25.7	7	10 Q49223	Q49223 glycine max
4	27	25.7	18	8 Q8KY0	Q8KY0 tuscutea ref
5	26	24.8	15	5 Q8TXC8	Q8TXC8 locusta mig
6	26	24.8	18	2 Q9R4N5	Q9R4N5 bacillus an
7	25	23.8	17	15 Q9DRS5	Q9DRS5 human immun
8	24.5	23.3	16	13 Q9PRY2	Q9PRY2 petromyzon
9	24	22.9	9	15 Q12096	Q12096 caprine art
10	24	22.9	9	15 Q12100	Q12100 caprine art
11	24	22.9	9	15 Q12102	Q12102 caprine art
12	24	22.9	9	15 Q12098	Q12098 caprine art
13	24	22.9	11	11 Q12104	Q12104 caprine art
14	24	22.9	11	11 Q99UC3	Q99UC3 rattus sp.
15	24	22.9	12	15 Q12090	Q12090 caprine art
16	24	22.9	12	15 Q12094	Q12094 caprine art

17	24	22.9	12	15	Q12114	O12114 caprine art
18	24	22.9	12	15	Q12082	Q12082 caprine art
19	24	22.9	12	15	Q12106	Q12106 caprine art
20	24	22.9	12	15	Q12092	Q12092 caprine art
21	24	22.9	12	15	Q12108	Q12108 caprine art
22	24	22.9	12	15	Q12074	Q12074 caprine art
23	24	22.9	12	15	Q12116	Q12116 caprine art
24	24	22.9	12	15	Q12118	Q12118 caprine art
25	24	22.9	12	15	Q12110	Q12110 caprine art
26	24	22.9	12	15	Q12112	Q12112 caprine art
27	24	22.9	12	15	Q12076	Q12076 caprine art
28	24	22.9	12	15	Q12088	Q12088 caprine art
29	24	22.9	12	15	Q12078	Q12078 caprine art
30	24	22.9	12	15	Q12080	Q12080 caprine art
31	24	22.9	12	15	Q12084	Q12084 caprine art
32	24	22.9	12	15	Q12086	Q12086 caprine art
33	24	22.9	15	2	Q9R531	Q9R531 thermus. ch
34	24	22.9	17	12	Q85004	Q85004 porcine res
35	23	21.9	9	8	Q94XE6	Q94XE6 tectocoris
36	23	21.9	10	2	Q8KH9	Q8KH9 clostridium
37	23	21.9	13	8	Q99783	Q99783 caprimulgus
38	23	21.9	14	4	Q13022	Q13022 homo sapien
39	23	21.9	15	4	Q9UC67	Q9UC67 homo sapien
40	23	21.9	15	8	Q8SJ19	Q8SJ19 phalacrocor
41	23	21.9	16	4	Q9UC54	Q9UC54 homo sapien
42	23	21.9	16	6	Q95MB4	Q95MB4 equus cabal
43	23	21.9	16	11	Q80WI5	Q80WI5 mus sp. ago
44	23	21.9	17	8	Q85UD6	Q85UD6 conger myri
45	23	21.9	17	11	P97758	P97758 mus musculu

ALIGNMENTS

RESULT 1

O14379 PRELIMINARY; PRT; 15 AA.
AC O14379
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Hypothetical protein (Fragment).
OS Schizosaccharomyces pombe (fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=972h-;
RA Jang Y.-J., Yoo H.-S.;
RL Submitted (APR-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; U97375; AAB63867.1; -.
KW Hypothetical protein.
FT NON_TER
SQ SEQUENCE 15 AA; 1636 MW; 93D127B36BBAF110 CRC64;

Query Match 32.4%; Score 34; DB 3; Length 15;
Best Local Similarity 54.5%; Pred. No. 1.3e+02;
Matches 6; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 5 VEWGCDXGCR 15
Db 4 VDYGCMSLSCR 14

RESULT 2

Q09053 PRELIMINARY; PRT; 18 AA.
AC Q09053
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)

DE Methanol dehydrogenase subunit 2 (EC 1.1.99.8) (MDH small beta
DE subunit) (Fragment).
GN MOXI.
OS Methylobacterium SP.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Methylococcales;
OC Methylococcaceae; Methylobacterium.
OX NCBI_TaxID=418;
RN [1]
RP SEQUENCE.
RC STRAIN=A4;
RX MEDLINE=93285990; PubMed=7685335;
RA Waechter-Brullia D., Dispirito A.A., Chistoserdova L.V., Lidstrom M.E.;
RL J. Bacteriol. 175:3767-3775(1993).
CC -!- CATALYTIC ACTIVITY: PRIMARY ALCOHOL + ACCEPTOR = ALDEHYDE +
CC REDUCED ACCEPTOR.
CC -!- COFACTOR: POO.
CC -!- SUBUNIT: THE HOLOENZYME MDH HAS AN ALPHA-2/BETA-2 CONFIGURATION OF
CC -!- SUBUNIT ALPHA AND A SMALL BETA SUBUNIT.
CC -!- SUBCELLULAR LOCATION: PERIPLASMIC.
DR GO:0019468; F:alcohol dehydrogenase (acceptor) activity; IEA.
DR GO:0004022; F:alcohol dehydrogenase activity; IEA.
DR GO:0016491; F:oxidoreductase activity; IEA.
DR GO:0015946; P:methanol oxidation; IEA.
DR InterPro: IPR003420; Meth_dh_beta.
DR Pfam: PF02315; MDH; 1.
DR Oxidoreductase; F00; Methanol utilization; Periplasmic.
KW NON_TER 1
FT NON_TER 18
FT NON_TER 18
SQ SEQUENCE 18 AA; 2014 MW; 2B0412D8D2B8B52D CRC64;

Query Match 28.6%; Score 30; DB 2; Length 18;
Best Local Similarity 50.0%; Pred. No. 6.7e+02;
Matches 4; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 10 DSXGCRPG 17
DB 2 DGTNCKPG 9

RESULT 3
ID O49223 PRELIMINARY; PRT; 7 AA.
AC O49223;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE HMG-1-like protein (Fragment).
OS Glycine max (Soybean).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosoids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae; Glycine.
OX NCBI_TaxID=3847;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Essex; TISSUE=Root;
RX MEDLINE=91367679; PubMed=1891369;
RA Laux T., Goldberg R.B.;
RL Submitted (FEB-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF047050; AAC03556.1; -.
FT NON_TER 1
FT NON_TER 1
SQ SEQUENCE 7 AA; 850 MW; 6AAAAAB378637810 CRC64;

Query Match 25.7%; Score 27; DB 10; Length 7;
Best Local Similarity 80.0%; Pred. No. 1e+06;
Matches 4; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 GWGCD 10
DB 1 GWGWD 5

RESULT 4
ID Q8SKY0 PRELIMINARY; PRT; 18 AA.
AC Q8SKY0;
DT 01-JUN-2002 (TrEMBLrel. 21, Created)
DT 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Ribosomal protein S11 (Fragment).
GN RPS11.
OS Cuscuta reflexa (Southern Asian dodder).
OC Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; asterids;
OC Lamiales; Solanales; Convolvulaceae; Cuscuta.
OX NCBI_TaxID=4129;
RN [1]
RP SEQUENCE FROM N.A.
RA Berg S.;
RT "Sequence analysis and coding potential of the holoparasitic flowering
RT plant genus Cuscuta";
RL Thesis (2002), Department of Institute of Botany, .
DR EMBL; AJ439611; CAD28796.1; -.
DR GO:0009507; C:chloroplast; IEA.
DR GO:0005622; C:intracellular; IEA.
DR GO:0005840; C:ribosome; IEA.
DR GO:0003735; F:structural constituent of ribosome; IEA.
DR GO:0006412; P:protein biosynthesis; IEA.
DR InterPro: IPR001971; Ribosomal_S11.
DR Pfam: PF00411; Ribosomal_S11; 1.
KW Chloroplast.
FT NON_TER 1
FT NON_TER 1
SQ SEQUENCE 18 AA; 2088 MW; 130D427BFE680B24 CRC64;

Query Match 25.7%; Score 27; DB 8; Length 18;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 GCRP 16
DB 9 GCRP 12

RESULT 5
ID Q9TXC8 PRELIMINARY; PRT; 15 AA.
AC Q9TXC8;
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE PROPHENOXIDASE inhibitor N terminus (Fragment).
OS Locusta migratoria (Migratory locust).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Orthopteroidea; Orthoptera; Caelifera; Acridoidea;
OC Acridoidea; Acrididae; Oedipodinae; Locusta.
OX NCBI_TaxID=7004;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91379003; PubMed=1910340;
RA Brehelin M., Boigegrain R.A., Drif L., Coletti-Previero M.A.;
RT "Purification of a protease inhibitor which controls prophenoloxidase
RT activation in hemolymph of Locusta migratoria (insecta).";
RL Biochem. Biophys. Res. Commun. 179:841-846(1991).
DR InterPro: IPR009041; PMP_inhibitor.
FT NON_TER 1
FT NON_TER 15
FT NON_TER 15
SQ SEQUENCE 15 AA; 1707 MW; C690EAE0112166C7 CRC64;

Query Match 24.8%; Score 26; DB 5; Length 15;

Best Local Similarity 80.0%; Pred. No. 2.3e+03;
Matches 4; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 14 CRPQ 18
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|
|
|
4 CTPQ 8

Db

RESULT 6

Q9R4N5 PRELIMINARY; PRT; 18 AA.

AC Q9R4N5; (Created)

DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)

DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)

DE EXTRACHTABLE antigen 1 (Fragment).

OS Bacillus anthracis.

OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.

OX NCBI_TaxID=1392;

RN [1]

RP MEDLINE=95247684; PubMed=7730281;

RA Farach J.W., Ribot W.J., Downs M.B., Ezzell J.W.;

RT "Purification and characterization of the major surface array protein

RL J. Bacteriol. 177:2481-2489 (1995).

SQ SEQUENCE 18 AA; 1926 MW; 1DBEFOA4925EFB6 CRC64;

Query Match 24.8%; Score 26; DB 2; Length 18;
Best Local Similarity 80.0%; Pred. No. 2.8e+03;
Matches 4; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 7 WGCDS 11
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|
|
13 WGCDS 17

Db

RESULT 7

Q9DRS5 PRELIMINARY; PRT; 17 AA.

AC Q9DRS5;

DT 01-MAR-2001 (TrEMBLrel. 16, Created)

DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)

DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)

DE Truncated nef protein (Negative factor) (27 kDa protein).

GN NEF.

OS Human immunodeficiency virus 1.

OC Viruses; Retroviridae; Retroviridae; Lentivirus.

OX NCBI_TaxID=11676;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=LTS 25e;

RA Aston L., Rhodes D., Solomon A., Deacon N., Satchell C., Carr A.,

RA Cooper D., Bitt R., Stewart G., Kaldor J.;

RT "Viral diversity in the nef/LTR region of the HIV-1 genome:

RL associated with long-term nonprogression.";

RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.

CC -!- FUNCTION: NEF HAS GTPASE, GTP-BINDING AND AUTOPHOSPHORYLATING

CC ACTIVITIES. IT SEEMS TO DOWN-REGULATE THE CD4(T4) ANTIGEN (BY

CC SIMILARITY).

DR EMBL; AF219708; AAG44185.1; --

DR GO; GO:0005525; P:GTP binding; IEA.

DR InterPro; IPR001558; HIV_Nef.

DR Pfam; PF00469; F-protein; 1.

DR AIDS; GTP-binding; Lipoprotein; Myristate.

SQ SEQUENCE 17 AA; 1846 MW; 656A7A36FEB808E CRC64;

Query Match 23.8%; Score 25; DB 15; Length 17;
Best Local Similarity 28.6%; Pred. No. 3.8e+03;
Matches 2; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 QDTIVGW 7
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|
|
|

Db 7 KSSVIGW 13

RESULT 8

Q9PRY2 PRELIMINARY; PRT; 16 AA.

AC Q9PRY2;

DT 01-MAY-2000 (TrEMBLrel. 13, Created)

DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)

DE Lectin 30 kDa subunit (fragment).

OS Petromyzon marinus (Sea lamprey).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Hyperoartia;

OC Petromyzontiformes; Petromyzontidae; Petromyzon.

OX NCBI_TaxID=7757;

RN [1]

RP SEQUENCE.

RX MEDLINE=94249896; PubMed=8192354;

RA Schluter S.F., Schroeder J., Wang E., Marchalonis J.J.;

RT "Recognition molecules and immunoglobulin domains in invertebrates.";

RL Ann. N. Y. Acad. Sci. 712:74-81 (1994).

SQ SEQUENCE 16 AA; 1728 MW; 3BBF03DD4185F446 CRC64;

Query Match 23.3%; Score 24.5; DB 13; Length 16;
Best Local Similarity 50.0%; Pred. No. 4.3e+03;
Matches 4; Conservative 1; Mismatches 2; Indels 1; Gaps 1;

QY 7 WGCDSXGC 14
|
|
|
|
1 WSC-TKGC 7

Db

RESULT 9

O12096 PRELIMINARY; PRT; 9 AA.

AC O12096;

DT 01-JUL-1997 (TrEMBLrel. 04, Created)

DT 01-JUL-1997 (TrEMBLrel. 04, Last sequence update)

DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)

DE Tat protein (fragment).

GN TAT.

OS Caprine arthritis encephalitis virus (CAEV).

OC Viruses; Retroviridae; Retroviridae; Lentivirus.

OX NCBI_TaxID=11660;

RN [1]

RP SEQUENCE FROM N.A.

RA Turelli P., Guiguen F., Mornex J.-F., Vigne R., Querat G.;

RT "dUTPase minus CAEV is attenuated for pathogenesis and accumulates G

RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.

DR EMBL; U81439; AA560832.1; --

FT NON_TER 1

SQ SEQUENCE 9 AA; 922 MW; 21E8644EB7340EB8 CRC64;

Query Match 22.9%; Score 24; DB 15; Length 9;
Best Local Similarity 44.4%; Pred. No. 1e+06;
Matches 4; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 9 CDSXGCRPG 17
|
|
|
|
1 CGCRLCNPG 9

Db

RESULT 10

O12100 PRELIMINARY; PRT; 9 AA.

AC O12100;

DT 01-JUL-1997 (TrEMBLrel. 04, Created)

DT 01-JUL-1997 (TrEMBLrel. 04, Last sequence update)

DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)

DE Tat protein (fragment).

GN TAT.

OS Caprine arthritis encephalitis virus (CAEV).

OC Viruses; Retrovirus; Retroviridae; Lentivirus.

OX NCBI_TaxID=11660;
RN [1]
RA Turelli P., Guiguen F., Mornex J.-F., Vigne R., Querat G.;
RP "dUTPase minus CAEV is attenuated for pathogenesis and accumulates G
RT to A substitutions.";
RT Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; U81441; AAB60836.1; -.
FT NON_TER 1
SQ SEQUENCE 9 AA; 922 MW; 21E8644EB7340EB8 CRC64;

Query Match 22.9%; Score 24; DB 15; Length 9;
Best Local Similarity 44.4%; Pred. No. 1e+06; 5; Indels 0; Gaps 0;
Matches 4; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 9 CDSXGCRPG 17
DB 1 CGCRLCNP 9

RESULT 11

O12102 PRELIMINARY; PRT; 9 AA.
AC O12102;
DT 01-JUL-1997 (TREMELrel. 04, Created)
DT 01-JUL-1997 (TREMELrel. 04, Last sequence update)
DT 01-DEC-2001 (TREMELrel. 19, Last annotation update)
DE Tat protein (fragment).
GN TAT.
OS Caprine arthritis encephalitis virus (CAEV).
OC Viruses; Retrovirus; Retroviridae; Lentivirus.
OX NCBI_TaxID=11660;
RN [1]
RA Turelli P., Guiguen F., Mornex J.-F., Vigne R., Querat G.;
RP "dUTPase minus CAEV is attenuated for pathogenesis and accumulates G
RT to A substitutions.";
RT Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; U81442; AAB60838.1; -.
FT NON_TER 1
SQ SEQUENCE 9 AA; 922 MW; 21E8644EB7340EB8 CRC64;

Query Match 22.9%; Score 24; DB 15; Length 9;
Best Local Similarity 44.4%; Pred. No. 1e+06; 5; Indels 0; Gaps 0;
Matches 4; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 9 CDSXGCRPG 17
DB 1 CGCRLCNP 9

RESULT 12

O12098 PRELIMINARY; PRT; 9 AA.
AC O12098;
DT 01-JUL-1997 (TREMELrel. 04, Created)
DT 01-JUL-1997 (TREMELrel. 04, Last sequence update)
DT 01-DEC-2001 (TREMELrel. 19, Last annotation update)
DE Tat protein (fragment).
GN TAT.
OS Caprine arthritis encephalitis virus (CAEV).
OC Viruses; Retrovirus; Retroviridae; Lentivirus.
OX NCBI_TaxID=11660;
RN [1]
RA Turelli P., Guiguen F., Mornex J.-F., Vigne R., Querat G.;
RP "dUTPase minus CAEV is attenuated for pathogenesis and accumulates G
RT to A substitutions.";
RT Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; U81440; AAB60835.1; -.
FT NON_TER 1
SQ SEQUENCE 9 AA; 922 MW; 21E8644EB7340EB8 CRC64;

Query Match 22.9%; Score 24; DB 15; Length 9;
Best Local Similarity 44.4%; Pred. No. 1e+06; 5; Indels 0; Gaps 0;
Matches 4; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 9 CDSXGCRPG 17
DB 1 CGCRLCNP 9

RESULT 13

O12104 PRELIMINARY; PRT; 9 AA.
AC O12104;
DT 01-JUL-1997 (TREMELrel. 04, Created)
DT 01-JUL-1997 (TREMELrel. 04, Last sequence update)
DT 01-DEC-2001 (TREMELrel. 19, Last annotation update)
DE Tat protein (fragment).
GN TAT.
OS Caprine arthritis encephalitis virus (CAEV).
OC Viruses; Retrovirus; Retroviridae; Lentivirus.
OX NCBI_TaxID=11660;
RN [1]
RA Turelli P., Guiguen F., Mornex J.-F., Vigne R., Querat G.;
RP "dUTPase minus CAEV is attenuated for pathogenesis and accumulates G
RT to A substitutions.";
RT Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; U81443; AAB60840.1; -.
FT NON_TER 1
SQ SEQUENCE 9 AA; 922 MW; 21E8644EB7340EB8 CRC64;

Query Match 22.9%; Score 24; DB 15; Length 9;
Best Local Similarity 44.4%; Pred. No. 1e+06; 5; Indels 0; Gaps 0;
Matches 4; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 9 CDSXGCRPG 17
DB 1 CGCRLCNP 9

RESULT 14

Q99JUC3 PRELIMINARY; PRT; 11 AA.
AC Q99JUC3;
DT 01-JUN-2001 (TREMELrel. 17, Created)
DT 01-JUN-2001 (TREMELrel. 17, Last sequence update)
DT 01-JUN-2003 (TREMELrel. 24, Last annotation update)
DE Luteinizing hormone/chorionic gonadotropin receptor homolog
DE (fragment).
OS Rattus sp.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10118;
RN [1]
RA Turelli P., Guiguen F., Mornex J.-F., Vigne R., Querat G.;
RP "dUTPase minus CAEV is attenuated for pathogenesis and accumulates G
RT to A substitutions.";
RT Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; U81440; AAB60835.1; -.
FT NON_TER 1
SQ SEQUENCE 11 AA; 994 MW; 333DCB137EB865B8 CRC64;

Query Match 22.9%; Score 24; DB 11; Length 11;
Best Local Similarity 50.0%; Pred. No. 3.5e+03; 1; Mismatches 3; Indels 0; Gaps 0;
Matches 4; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

OY 6 GWGCDXSG 13
Db 4 GSGCGAAG 11

RESULT 15

O12090
ID O12090 PRELIMINARY; PRT; 12 AA.
AC O12090;
DT 01-JUL-1997 (TREMELrel. 04, Created)
DT 01-JUL-1997 (TREMELrel. 04, Last sequence update)
DT 01-JUN-2003 (TREMELrel. 24, Last annotation update)
DE Tat protein (Fragment).
GN TAT.
OS Caprine arthritis encephalitis virus (CAEV).
OC Viruses; Retroviral viruses; Retroviridae; Lentivirus.
OX NCBI_TaxID=11660;
RN [1]
RP SEQUENCE FROM N.A.
RA Turelli P., Guiguen P., Mornex J.-F., Vigne R., Querat G.;
RT "dUTPase minus CAEV is attenuated for pathogenesis and accumulates G
to A substitutions.";
RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; U81436; A860826.1; -
DR GO; GO:0016563; P:transcriptional activator activity; IEA.
DR GO; GO:0045941; P:positive regulation of transcription; IEA.
DR InterPro; IPR004247; Lentiviral_Tat.
DR Pfam; PF02998; Lentiviral_Tat; I.
FT NON_TER 1
SQ SEQUENCE 12 AA; 1266 MW; 5A60BBB1E8644EB7 CRC64;

Query Match 22.9%; Score 24; DB 15; Length 12;
Best Local Similarity 44.4%; Pred. No. 3.8e+03;
Matches 4; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 9 CDSXGCRFG 17
Db 1 CGCRLCNFG 9

Search completed: March 13, 2004, 07:29:49
Job time : 42 secs